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3921 63985

Date 4/8 Serial # 091546127 Priority Application Date _____

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What is the topic, such as the novelty, motivation, utility, or other specific facets defining the
desired focus of this search? Please include the concepts, synonyms, keywords, acronyms,
registry numbers, definitions, structures, strategies, and anything else that helps to describe the
topic. Please attach a copy of the abstract and pertinent claims.

Claims 1-17 + 21-40
Problem See Page 1 Lines 14-31
Solution " " 9 " 1-9

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archer: <u>Derrick Blalock</u>		Structure (#) _____	STN <input checked="" type="checkbox"/>
archer Phone: _____		Bibliographic <input checked="" type="checkbox"/>	Dialog <input checked="" type="checkbox"/>
archer Location: STIC-EIC2800, CP4-9C18		Litigation <input type="checkbox"/>	Questel/Orbit _____
Searcher Picked Up: <u>4/8/02</u>		Fulltext <input type="checkbox"/>	Lexis-Nexis _____
Completed: <u>60 4/9/02</u>		Patent Family <input type="checkbox"/>	WWW/Internet _____
archer Prep/Rev Time: <u>60</u>		Other <input type="checkbox"/>	Other _____
line Time: <u>170</u>			

04/09/2002

Serial No.:09/846,127

L1 FILE 'REGISTRY' ENTERED AT 11:49:52 ON 09 APR 2002
139 S (TA AND O)/ELS AND 2/ELC.SUB

L2 FILE 'HCAPLUS' ENTERED AT 11:50:47 ON 09 APR 2002
11477 S L1
L3 36 S L2 AND (EMITTER OR ECL OR (COLLECTOR(2N) ELECTRODE))

L3 ANSWER 1 OF 36 HCAPLUS COPYRIGHT 2002 ACS
AN 2002:486 HCAPLUS
DN 136:206800
TI Anodic-oxidation growth of microscopic pillar arrays: kinetic aspects
AU Vorob'eva, A. I.
CS Belarussian State University of Information Science and Electronics,
Belarus
SO Russian Microelectronics (Translation of Mikroelektronika) (2001), 30(6),
381-393
CODEN: RUICE5; ISSN: 1063-7397
PB MAIK Nauka/Interperiodica Publishing
DT Journal
LA English
AB The three-step fabrication of microscopic pillar arrays by the anodic
oxidn. of Al/Ta thin-film structures on dielec. or silicon substrates was
studied exptl. The major features of pillar-growth kinetics are
described. The main properties of the arrays are evaluated by SEM and
simultaneous current-voltage tracing. The ranges of variation for
geometric array parameters are detd. The pillars grown have a max.
height-to-diam. ratio of 17.0, a max. height of 540 nm, and a min. radius
of .apprx.15 nm. The max. d. of pillars in an array is 8.25 .times. 1010
cm-2. A good reproducibility of phys. and morphol. properties is achieved
for large-area pillar arrays. Potential applications of pillar arrays are
recited: light-emitting diodes, thin-film controllers, solar batteries,
spatial light modulators, polarizers, etc. A study into the fabrication
of pillar arrays for field-emitter displays is currently in
progress.
IT Breakdown voltage
(in anodization of Al/Ta thin film structures in growth of microscopic
pillar arrays)
IT 7429-90-5, Aluminum, uses 7440-25-7, Tantalum, uses
RL: CPS (Chemical process); DEV (Device component use); PEP (Physical,
engineering or chemical process); PRP (Properties); PROC (Process); USES
(Uses)
(anodization of Al/Ta thin film structures in growth of microscopic
pillar arrays: kinetic aspects)
IT 7440-21-3, Silicon, uses
RL: DEV (Device component use); PRP (Properties); USES (Uses)
(anodization of Al/Ta thin film structures on silicon in growth of
microscopic pillar arrays)
IT 144-62-7, Oxalic acid, uses 7664-38-2, Phosphoric acid, uses
7664-93-9, Sulfuric acid, uses

L3 ANSWER 2 OF 36 HCAPLUS COPYRIGHT 2002 ACS
AN 2001:907177 HCAPLUS
DN 136:12566
TI Fiber-optic Raman lasers - design versions
IN Dianov, E. M.; Bufetov, I. A.; Grekov, M. V.; Karpov, V. I.; Prokhorov, A.
M.
PA Nauchnyi Tsentr Volokonnoi Optiki pri Institute Obshchel Fiziki RAN,
Russia
SO Russ., No pp. given
CODEN: RUXXE7
DT Patent
LA Russian
FAN.CNT 1
PATENT NO. KIND DATE APPLICATION NO. DATE

PI RU 2152676 C1 20000710 RU 1998-117721 19980922
 AB Raman fiber lasers are described which comprise a length of a phosphate fiber contg. .gtoreq.1 addnl. element as an active medium; a pumping source; and optical elements allowing multiple passing through the fiber waveguide portion of Stokes components bonded with phosphor oxide; and which contains a long-period grating ensuring addnl. optical loss in the Stokes component bonded with the .gtoreq.1 addnl. element. Near-IR Raman fiber lasers in which the pumping lasers have yttrium ions, neodymium ions or chromium (4+) ions as **emitters** were discussed.

IT (variations in design of phosphate fiber Raman laser)
 15118-03-3, Forsterite 50814-00-1, Calcium germanium oxide
 RL: DEV (Device component use); USES (Uses)
 (chromium-doped; variations in design of phosphate fiber Raman laser contg.)
 IT 1303-86-2, Boron oxide, occurrence 1304-28-5, Barium oxide, occurrence 1304-76-3, Bismuth oxide, occurrence 1314-23-4, Zirconium oxide, occurrence 1332-29-2, Tin oxide 1332-37-2, Iron oxide, occurrence 1344-28-1, Aluminum oxide, occurrence 7631-86-9, Silicon oxide, occurrence 7783-41-7, Fluorine oxide 11104-93-1, Nitrogen oxide, occurrence 12024-21-4, Gallium oxide 13463-67-7, Titanium oxide, occurrence 59763-75-6, Tantalum oxide
 RL: DEV (Device component use); OCU (Occurrence, unclassified); OCCU (Occurrence); USES (Uses)
 (phosphate fiber contg.; variations in design of phosphate fiber Raman laser contg.)
 IT 7440-00-8, Neodymium, uses 7440-47-3, Chromium, uses 7440-64-4, Ytterbium, uses
 RL: DEV (Device component use); MOA (Modifier or additive use); USES (Uses)

L3 ANSWER 3 OF 36 HCAPLUS COPYRIGHT 2002 ACS
 AN 2000:823090 HCAPLUS
 DN 133:368551
 TI Field-emitting electron source
 IN Yamakishi, Toshio; Nanba, Masakazu; Okazaki, Saburo; Hirano, Yoshiyuki; Okamura, Noritomo; Katsuhara, Yukinori; Inoue, Shigeru
 PA Japan Broadcasting Corp., Japan; Hitachi Electronics Co., Ltd.
 SO Jpn. Kokai Tokkyo Koho, 11 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI JP 2000323011 A2 20001124 JP 1999-129122 19990510
 AB The electron source, from which electrons are emitted by applying elec. voltage on a cathode and gate electrodes facing each other and the space sandwiched between the electrodes involves a porous elec. insulator. The elec. insulator has fine pores extended in the thickness direction as a result of anodization and pores involve **emitters**. Alternatively, the pores in the elec. insulator are formed by etching through a mask made of an anodized porous film having fine pores in the direction perpendicular to the thickness direction. The electron source with having submicron- to nano-order **emitters** can be obtained without photolithog., i.e., at low cost.
 IT Anodization
 Electric insulators

Etching

Field emitters

(field-emitting electron source having elec. insulator involving
emitter in micropores formed by anodization or etching)

L3 ANSWER 4 OF 36 HCAPLUS COPYRIGHT 2002 ACS
 AN 2000:665686 HCAPLUS
 DN 133:246254
 TI Electron-emitting material with low evaporation during discharge and
resistant to ion sputtering and its low-cost fabrication
 IN Hamada, Munemitsu; Takeishi, Akira; Takahashi, Makoto; Matsuoka, Dai;
Yodogawa, Masatada; Harada, Hiraku
 PA TDK Corporation, Japan
 SO Eur. Pat. Appl., 37 pp.
 CODEN: EPXXDW
 DT Patent
 LA English
 FAN.CNT 2

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 1037244	A2	20000920	EP 2000-301668	20000301
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
JP 3078287	B1	20000821	JP 1999-346966	19991206
JP 2000331603	A2	20001130		

AB An electron-emitting material contains a 1st metal component selected from Ba, Sr and Ca and a 2nd metal component selected from Ta, Zr, Nb, Ti and Hf and also contains oxynitride perovskite. The electron-emitting material has improved electron emission characteristics, restrained evapn. at elevated temps., and minimized consumption by ion sputtering. The electron-emitting material was prep'd. by firing a metal component-contg. raw material disposed in proximity to C in a N gas-contg. atm. to thereby create oxynitride perovskite.

IT Cathodes
 Electrodes
 Field emission cathodes
 Photocathodes
 Sputtering cathodes
 (fabrication of electron emitters)

L3 ANSWER 5 OF 36 HCAPLUS COPYRIGHT 2002 ACS
 AN 2000:574048 HCAPLUS
 DN 133:153231
 TI Antireflection coated refractory metal matched IR emitter for
use in thermophotovoltaic generators
 IN Fraas, Lewis M.; Magendanz, Galen; Avery, James E.
 PA Jx Crystals Inc., USA
 SO PCT Int. Appl., 21 pp.
 CODEN: PIXXD2
 DT Patent
 LA English
 FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2000048231	A2	20000817	WO 1999-US24736	19991022
W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN,				

WO 1999-US24736 W 19991022

AB Thermophotovoltaic (TPV) elec. power generators have **emitters** with IR outputs matched with usable wavelengths for converter cells. The **emitters** have durable substrates, optional refractory isolating layers, conductive refractory metal or inter-metallic **emitter** layers, and refractory metal oxide antireflection layers. SiC substrates have W or TaSi₂ **emitter** layers and 0.14 .mu. ZrO₂ or Al₂O₃ antireflection layers used as IR **emitters** for GaSb converter cells in TPV generators.

IT 409-21-2, Silicon carbide (SiC), uses 1314-23-4, Zirconium oxide (ZrO₂), uses 1314-61-0, Tantalum pentoxide 1344-28-1, Alumina, uses 7439-98-7, Molybdenum, uses 7440-03-1, Niobium, uses 7440-25-7, Tantalum, uses 7440-33-7, Tungsten, uses 12034-80-9, Niobium disilicide 12039-79-1, Tantalum silicide (TaSi₂) 12039-83-7, Titanium disilicide 12039-87-1, Vanadium disilicide 12064-03-8 12597-68-1, Stainless steel, uses 12611-97-1, Kanthal 273921-59-8, Nichrome
 RL: DEV (Device component use); USES (Uses)
 (antireflection coated refractory metal matched IR **emitter** for use in thermophotovoltaic generators)

L3 ANSWER 6 OF 36 HCAPLUS COPYRIGHT 2002 ACS

AN 2000:541583 HCAPLUS

DN 133:289670

TI Studies on the interaction between thin film materials and Mo field **emitter** arrays

AU Chalamala, Babu R.; Reuss, Robert H.

CS Flat Panel Display Division, Motorola Incorporated, Tempe, AZ, 85284, USA

SO Journal of Vacuum Science & Technology, B: Microelectronics and Nanometer Structures (2000), 18(4), 1825-1832

CODEN: JVTBD9; ISSN: 0734-211X

PB American Institute of Physics

DT Journal

LA English

AB A simple method for the evaluation of materials suitable for the fabrication of field emission vacuum microelectronic devices is presented. Since there can be a wide range of electron and ion interactions with the device, it is important to be able to quickly assess if a material may have a particular adverse effect on emission performance under operational conditions. The technique is based on the sensitivity of a large field **emitter** array to the outgassing or desorption of gas species from thin films under electron beam excitation. Mo field **emitter** arrays degraded rapidly with stainless steel anodes coated with various oxide materials. The extent of degrdn. is the most rapid with SiO₂, Si₃N₄, and MoO₃ thin films. Stainless steel anodes with Mo and Nb thin films show a faster degrdn. rate than stainless steel anodes, most likely because of native oxides grown during processing and handling. The emission behavior in the presence of Ir, Pd, Al, Zn, and Ti metal films and barrier materials like C and TaN is similar to stainless steel ref. data. Once the oxide films are covered with barrier layers like C and TaN, emission decay rates approach the values obtained with stainless steel ref. anodes. The obsd. emission current degrdn. is consistent with a model based on the liberation of oxygen from the surface of electron beam bombarded materials. Using controlled oxygen exposure expts., the authors detd. the equiv. local oxygen pressures in the presence of various thin films. With thin films of Nb, ZrO₂, Ta₂O₅, MgO, Nb₂O₅, and Al₂O₃, the emission degrdn. is akin to having a local O₂ partial pressure in the 1 .times. 10⁻⁷-1 .times. 10⁻⁶ Torr range and with Mo, MoO₃, Si₃N₄, and SiO₂, this is equiv. to having local O₂ pressures of 1 .times. 10⁻⁵ Torr.

IT Electron beams

Field emission
 Field emitters
 Microelectronic devices
 (studies on the interaction between thin film materials and Mo field
 emitter arrays)
 IT 7439-98-7, Molybdenum, properties
 RL: DEV (Device component use); PRP (Properties); TEM (Technical or
 engineered material use); USES (Uses)
 (studies on the interaction between thin film materials and Mo field
 emitter arrays)
 IT 1309-48-4, Magnesium oxide (MgO), properties 1313-27-5, Molybdenum oxide
 (MoO₃), properties 1313-96-8, Niobium oxide (Nb₂O₅) 1314-23-4,
 Zirconium oxide (ZrO₂), properties 1314-61-0, Tantalum oxide
 (Ta₂O₅) 1344-28-1, Alumina, properties 7429-90-5, Aluminum, properties
 7439-88-5, Iridium, properties 7440-03-1, Niobium, properties
 7440-05-3, Palladium, properties 7440-32-6, Titanium, properties
 7440-66-6, Zinc, properties 7631-86-9, Silica, properties 12033-89-5,
 Silicon nitride (Si₃N₄), properties 12597-68-1, Stainless steel,
 properties
 RL: MOA (Modifier or additive use); PRP (Properties); RCT (Reactant); RACT
 (Reactant or reagent); USES (Uses)
 (studies on the interaction between thin film materials and Mo field
 emitter arrays)
 RE.CNT 20 THERE ARE 20 CITED REFERENCES AVAILABLE FOR THIS RECORD
 ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 7 OF 36 HCPLUS COPYRIGHT 2002 ACS
 AN 2000:474520 HCPLUS
 DN 133:96633
 TI Image display devices
 IN Mitamura, Satoshi
 PA Sony Corp., Japan
 SO Jpn. Kokai Tokkyo Koho, 11 pp.
 CODEN: JKXXAF
 DT Patent
 LA Japanese
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 2000195671	A2	20000714	JP 1998-368000	19981224
AB	The devices comprise: (1) a glass substrate; (2) a red, a green and a blue phosphor matrix; (3) an ITO 1st electrode stripe array (.dblvert. X); (4) a UV-emitting layer (lamine); (5) a dielec. layer; and (6) a 2nd electrode (.dblvert. Y), where (4) typically comprises AlN:Gd, or a semiconductor LED comprising a GaN/AlGaN-MQW laminate.				
IT	Fluorescent substances Optical imaging devices Quantum well devices Quantum well heterojunctions UV radiation (image display devices)				
IT	1314-61-0, Tantalum oxide (Ta ₂ O ₅) 1314-98-3, Zinc sulfide (ZnS), uses 7439-98-7, Molybdenum, uses 12340-04-4, Yttrium oxide sulfide (Y ₂ O ₂ S) 24304-00-5, Aluminum nitride (AlN) 50926-11-9, ITO 117656-36-7, Aluminum gallium nitride (Al _{0.3} Ga _{0.7} N) 120831-83-6, Aluminum gallium nitride (Al _{0.25} Ga _{0.75} N) 125297-45-2, Aluminum gallium nitride al0.2ga0.8n RL: DEV (Device component use); USES (Uses) (image display devices)				

IT 7429-90-5, Aluminum, uses 7440-22-4, Silver, uses 7440-27-9, Terbium, uses 7440-50-8, Copper, uses 7440-53-1, Europium, uses
 RL: MOA (Modifier or additive use); USES (Uses)
 (image display devices)

L3 ANSWER 8 OF 36 HCAPLUS COPYRIGHT 2002 ACS

AN 2000:470322 HCAPLUS

DN 133:98270

TI Fabrication of memory elements from composites of phase-change materials and dielectric materials

IN Ovshinsky, Standford R.; Czubatyj, Wolodymyr; Strand, David A.; Klersy, Patrick J.; Kostylev, Sergey; Pashmakov, Boil

PA Energy Conversion Devices, Inc., USA

SO U.S., 12 pp., 5825046Cont.-in-part of U.S. 5,825,046.

CODEN: USXXAM

DT Patent

LA English

FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 6087674	A	20000711	US 1998-63174	19980420
	US 5825046	A	19981020	US 1996-739080	19961028
	KR 2000052840	A	20000825	KR 1999-703674	19990427

PRAI US 1996-739080 A2 19961028

AB An elec. operated, single cell memory element comprising: a vol. of memory material defining a single-cell memory element, the memory material comprising a heterogeneous mixt. of a phase-change material and a dielec. material; and means for delivering an elec. signal to at least a portion of the vol. of memory material. An elec. operated, single-cell memory element comprising: a vol. of memory material defining the single-cell memory element, the memory material comprising a phase-change material and a dielec. material where the phase-change material has a plurality of detectable resistivity values and can be set directly to one of the resistivity values without the need to be set to a specific starting or erased resistivity value, regardless of the previous resistivity value of the material, in response to an elec. signal; and means for delivering the elec. signal to at least a portion of the vol. of memory material.

IT 1310-53-8, Germanium oxide (GeO₂), processes 1314-23-4, Zirconium oxide (ZrO₂), processes 1314-61-0, Tantalum oxide (Ta₂O₅) 1314-98-3, Zinc sulfide (ZnS), processes 7783-40-6, Magnesium fluoride (MgF₂) 7789-75-5, Calcium fluoride (CaF₂), processes 10043-11-5, Boron nitride, processes 12025-34-2, Germanium sulfide (GeS₂) 12033-89-5, Silicon nitride, processes 13759-10-9, Silicon sulfide (SiS₂) 24304-00-5, Aluminum nitride (AlN) 25583-20-4, Titanium nitride (TiN) 25658-42-8, Zirconium nitride (ZrN) 39327-44-1, Lithium fluoride (LiF₂)
 RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses)
 (dielec. materials; fabrication of memory elements from composites of phase-change materials and)

IT 12033-89-5D, Silicon nitride, non-stoichiometric
 RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses)
 (fabrication of memory elements from composites of phase-change materials and)

L3 ANSWER 9 OF 36 HCAPLUS COPYRIGHT 2002 ACS

AN 1999:398502 HCAPLUS

DN 131:109743

TI Doping effects of Groups III, IV, or V elements on the **emitter** of CRT oxide cathodes

AU Hayashida, Yoshiki; Ozawa, Tetsuro; Sakurai, Hiroshi

CS Matsushita Electronics Corporation, Osaka, 569-1193, Japan

SO Appl. Surf. Sci. (1999), 146(1-4), 7-11

CODEN: ASUSEE; ISSN: 0169-4332

PB Elsevier Science B.V.

DT Journal

LA English

AB Doping effects on CRT oxide cathodes was studied with dopant elements of groups III, IV, or V. Doping effects correspond neg. to both resistance of the **emitter** layer and m.ps. of dopants. The doping effect mechanism is discussed in terms of elec. cond. of the **emitter** layer with interaction between dopants and reducing agents. Based on the idea that diffusibility of dopants could improve cond., the highest doping effect was achieved with ZrO₂ copptn. doping.

IT Melting point
(dopant; doping effects of Groups III, IV, or V element oxides on the **emitter** of CRT barium strontium oxide cathodes)

IT 1308-96-9, Europium oxide (Eu₂O₃) 1313-96-8, Niobium oxide (Nb₂O₅)
1314-23-4, Zirconium oxide (ZrO₂), properties 1314-36-9, Yttrium oxide (Y₂O₃), properties 1314-61-0, Tantalum oxide (Ta₂O₅)
1314-62-1, Vanadium oxide (V₂O₅), properties 12037-01-3, Terbium oxide (Tb₄O₇) 12055-23-1, Hafnium oxide (HfO₂) 12060-08-1, Scandium oxide (Sc₂O₃) 13463-67-7, Titanium oxide (TiO₂), properties
RL: MOA (Modifier or additive use); PEP (Physical, engineering or chemical process); PRP (Properties); PROC (Process); USES (Uses)
(doping effects of Groups III, IV, or V element oxides on the **emitter** of CRT barium strontium oxide cathodes)

L3 ANSWER 10 OF 36 HCAPLUS COPYRIGHT 2002 ACS

AN 1999:296754 HCAPLUS

DN 130:319638

TI Electron **emitters** having an etching-resistant insulator

IN Nakanishi, Masayuki; Chin, Kazutami; Shimnojo, Norihide

PA Ise Electronics Corp., Japan

SO Jpn. Kokai Tokkyo Koho, 5 pp.
CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
-----	-----	-----	-----	-----
PI JP 11120897	A2	19990430	JP 1997-276950	19971009

AB The title **emitters** comprise pointy-projection **emitter** electrodes formed on a Si substrate, an insulator layer having openings to expose around the tops of the pointy electrodes, and gate electrodes each formed around the **emitters** on the insulator for impression of voltage across the **emitters** and the gate electrodes. The insulator layer is made from an etching resistant against HF and may be chosen from Al₂O₃, TiO₂, Ta₂O₅, or Si₃N₄. The use of the insulator material prevents over-etching of the insulator in otherwise causing short-circuiting the **emitter** and gate electrodes.

IT Cathodes
(electron **emitters** having an etching-resistant insulator)

L3 ANSWER 11 OF 36 HCAPLUS COPYRIGHT 2002 ACS

AN 1999:292322 HCAPLUS

DN 131:26128
 TI Development of the oxygenated thermionic energy converters utilizing the sputtered metal oxides as a collector
 AU Fukuda, Ryuzo; Kasuga, Yasuhiro; Kato, Ken; Shimizu, Sadaaki
 CS Electrotechnical Laboratory, Tsukuba, 305-8568, Japan
 SO AIP Conf. Proc. (1999), 458 (Space Technology and Applications International Forum--1999, Pt. 2), 1444-1451
 CODEN: APCPCS; ISSN: 0094-243X
 PB American Institute of Physics
 DT Journal; General Review
 LA English
 AB A review with 7 refs. Refractory metal oxides such as NbO_x, WO_x, MoO_x, TaO_x, PtO_x and Silver oxide (AgO_x) were studied for an oxygenated thermionic converter and a low work function collector. The metal oxide materials were deposited on metal substrates by RF sputtering in the Ar/O₂ gas mixt. to fabricate **collector electrodes**. Work function values of the **collector electrodes** were evaluated by cesium plasma immersion technique. Very low work function values were obtained, for instance 1.25 V as ϕ of AgO_x. A research thermionic energy converter with a plane parallel type of a polycryst. W **emitter** and the metal oxide collectors (AgO_x, NbO_x and PtO_x), was set up and the power generation expts. were conducted. In the case of the W-AgO_x thermionic converter, the barrier index VB was 2.05 V. Similarly VB = 2.15V for the W-NbO_x converter, VB = 2.25V for the W-PtO_x converter.
 IT Energy converters
 (thermonic; development of the oxygenated thermionic energy converters utilizing the sputtered metal oxides as a collector)
 IT 1314-35-8, Tungsten oxide (WO₃), properties 11098-99-0, Molybdenum oxide 11129-89-8, Platinum oxide 12627-00-8, Niobium oxide 20667-12-3, Silver oxide (Ag₂O) 59763-75-6, Tantalum oxide
 RL: PEP (Physical, engineering or chemical process); PRP (Properties); TEM (Technical or engineered material use); PROC (Process); USES (Uses)
 (development of the oxygenated thermionic energy converters utilizing the sputtered metal oxides as a collector)
 RE.CNT 10 THERE ARE 10 CITED REFERENCES AVAILABLE FOR THIS RECORD
 ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 12 OF 36 HCPLUS COPYRIGHT 2002 ACS
 AN 1998:719906 HCPLUS
 DN 130:44558
 TI Enhancement of Electrochemical Hot Electron Injection into Electrolyte Solutions at Oxide-Covered Tantalum Electrodes by Thin Platinum Films
 AU Sung, Yung-Eun; Bard, Allen J.
 CS Department of Chemistry and Biochemistry, The University of Texas at Austin, Austin, TX, 78712, USA
 SO J. Phys. Chem. B (1998), 102(49), 9806-9811
 CODEN: JPCBFK; ISSN: 1089-5647
 PB American Chemical Society
 DT Journal
 LA English
 AB The previously reported exptl. evidence for hot soln.-phase electrons generated at Ta₂O₅-covered Ta electrodes in both acetonitrile and aq. solns. using electrogenerated chemiluminescence (ECL) and electrochem. measurements was extended by observing the effect of thin Pt films. Hot electron injection was monitored by noting the photoemission following redn. of the thianthrene radical cation (TH.bul.+), signaling the direct formation of the excited state (TH.bul.+ + ehs .fwdarw. TH*), a process that does not occur at a bulk Pt electrode. We report the enhancement of hot electron injection efficiency by a factor of .apprx.5

by deposition of a thin (<40 nm) Pt film on the Ta/Ta₂O₅ electrode. This enhancement of ECL efficiency at the metal/oxide/Pt/liq. interface is ascribed to the suppression of the Ta₂O₅ surface states by the Pt film. The effect of Pt film thickness was investigated and showed a decrease in emission with increased film thickness, in accord with the expected mean free path of hot electrons in the Pt (ehPt). In examg. the Ta/Ta₂O₅/Pt/soln. system, one can contrast the behavior obsd. when an elec. connection is made directly to the Pt with that obsd. when a connection to the Pt is made via the Ta, where ehPt species are generated.

IT Electric potential
(effect on electrogenerated luminescence of thianthrene during electroredn. on Ta/Ta₂O₅ and Ta/Ta₂O₅/Pt electrodes)

IT Films
(formation, of tantalum pentoxide on tantalum and platinum on oxide covered tantalum)

IT X-ray photoelectron spectra
(of Ta/Ta₂O₅ and Ta/Ta₂O₅/Pt electrodes)

IT Surface structure
(of Ta/Ta₂O₅ and Ta/Ta₂O₅/Pt electrodes by AFM)

IT Thickness
(of platinum films on oxide covered tantalum electrodes, effect on electrochem hot electron injection into electrolyte soln)

IT Sputter deposition
(of platinum on oxide covered tantalum electrode)

IT Anodizing
(of tantalum in ammonium tartrate soln.)

IT Electrochemiluminescence
(of thianthrene radical cation following of redn. on oxide covered tantalum electrode in acetonitrile)

IT 3164-29-2, Ammonium tartrate
RL: NUU (Other use, unclassified); PRP (Properties); USES (Uses)
(anodization of tantalum in soln. of)

IT 7440-06-4, Platinum, properties
RL: FMU (Formation, unclassified); PRP (Properties); FORM (Formation, nonpreparative)
(deposition on oxide covered tantalum electrode by sputtering)

IT 7440-25-7, Tantalum, uses
RL: DEV (Device component use); PRP (Properties); USES (Uses)
(electrochem hot electron injection into electrolyte soln at oxide covered tantalum electrode by thin platinum film)

IT 183748-02-9, Electron
RL: PRP (Properties)
(electrochem hot electron injection into electrolyte soln. at oxide covered tantalum electrode by thin platinum film)

IT 583-52-8, Dipotassium oxalate
RL: NUU (Other use, unclassified); PRP (Properties); USES (Uses)
(electrogenerated luminescence of thianthrene during electroredn. on Ta/Ta₂O₅ and Ta/Ta₂O₅/Pt electrodes in soln. of)

IT 1314-61-0, Tantalum oxide Ta₂O₅
RL: FMU (Formation, unclassified); PRP (Properties); FORM (Formation, nonpreparative)
(formation on surface of tantalum by anodization in ammonium tartrate soln.)

L3 ANSWER 13 OF 36 HCPLUS COPYRIGHT 2002 ACS
AN 1998:719902 HCPLUS
DN 130:72880
TI Demonstration of Electrochemical Generation of Solution-Phase Hot Electrons at Oxide-Covered Tantalum Electrodes by Direct Electrogenerated

Chemiluminescence

AU Sung, Yung-Eun; Gaillard, Frederic; Bard, Allen J.
 CS Department of Chemistry and Biochemistry, The University of Texas at Austin, Austin, TX, 78712, USA
 SO J. Phys. Chem. B (1998), 102(49), 9797-9805
 CODEN: JPCBFK; ISSN: 1089-5647
 PB American Chemical Society
 DT Journal
 LA English
 AB Exptl. evidence for the prodn. of hot electrons in an acetonitrile soln. from a Ta205-covered Ta electrode was provided by electrogenerated chemiluminescence (**ECL**) and electrochem. measurements. Electron transfer to soln. species occurred via the Ta205 conduction band, as demonstrated by comparative measurements with a no. of 1-electron redox couples at Pt and Ta electrodes. The oxidized forms of thianthrene and a heptamethine cyanine dye were selected as the species capable of direct formation of the excited state and **ECL** upon hot electron injection. The observation of **ECL** emission upon a cathodic potential step (a process that does not occur at a metal electrode) confirmed the occurrence of this process. **ECL** emission at Ta/Ta205 was also obsd. during redn. of Ru(bpy)33+ (bpy = bipyridine). Reasons for the low efficiency of the **ECL** process via hot electrons at the metal/metal oxide/soln. interface are discussed.

IT Electrochemiluminescence
 Hot electrons
 (demonstration of electrochem. generation of soln.-phase hot electrons at oxide-covered tantalum electrodes by direct electrogenerated chemiluminescence)

IT 92-85-3, Thianthrene 69415-17-4
 RL: PRP (Properties); RCT (Reactant)
 (electrochem. oxidn.: demonstration of electrochem. generation of soln.-phase hot electrons at oxide-covered tantalum electrodes by direct electrogenerated chemiluminescence)

IT 91-15-6, Phthalonitrile 118-75-2, properties 129-00-0, Pyrene, properties 486-25-9, 9-Fluorenone 527-17-3, Duroquinone 1518-16-7, TCNQ 67994-95-0 88505-29-7, Bis(tetrabutylammonium) peroxydisulfate
 RL: PRP (Properties); RCT (Reactant)
 (electrochem. reactions at platinum and at Ta/Ta205: conduction band energies of Ta205 and electrochem. generation of soln.-phase hot electrons at oxide-covered tantalum electrodes by direct electrogenerated chemiluminescence)

IT 91-20-3, Naphthalene, properties
 RL: PRP (Properties); RCT (Reactant)
 (electrochem. reactions at platinum: conduction band energies of Ta205 and electrochem. generation of soln.-phase hot electrons at oxide-covered tantalum electrodes by direct electrogenerated chemiluminescence)

L3 ANSWER 14 OF 36 HCPLUS COPYRIGHT 2002 ACS
 AN 1998:196267 HCPLUS
 DN 128:264977
 TI Thin-film electron **emitter** for display apparatus
 IN Kusunoki, Toshiaki; Suzuki, Mutsumi
 PA Hitachi, Ltd., Japan
 SO Jpn. Kokai Tokkyo Koho, 6 pp.
 CODEN: JKXXAF
 DT Patent
 LA Japanese

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 10079221	A2	19980324	JP 1996-233914	19960904
AB	The invention relates to a thin-film MIM electron emitter for display app., wherein the use of a refractory metal lower electrode affords resistance to electromigration and stress migration.				
IT	Cathodes Optical imaging devices (MIM; thin-film electron emitter for display app.)				
IT	1314-61-0, Tantalum oxide	1344-28-1, Alumina, uses Aluminum, uses	7440-25-7, Tantalum, uses	7429-90-5,	RL: DEV (Device component use); USES (Uses)
	(thin-film electron emitter for display app.)				
L3	ANSWER 15 OF 36 HCPLUS COPYRIGHT 2002 ACS				
AN	1997:785859 HCPLUS				
DN	128:14896				
TI	Development of refractory metal oxide collector materials and their thermionic converter performance				
AU	Fukuda, R.; Kasuga, Y.; Katoh, K.				
CS	Energy Division, Electrotechnical Laboratory, Tsukuba, 305, Japan				
SO	Funct. Graded Mater. 1996, Proc. Int. Symp., 4th (1997), Meeting Date 1996, 647-654. Editor(s): Shiota, Ichiro; Miyamoto, Yoshinari. Publisher: Elsevier, Amsterdam, Neth.				
	CODEN: 65KZAW				
DT	Conference				
LA	English				
AB	Refractory metal oxides of NbO _x , WO _x , TaO _x , and AgO _x were studied for use as a high-performance collector. The metal oxide materials were deposited on metal substrates by radio-frequency sputtering in the Ar/O ₂ gas mixt., in which the partial pressure of O ₂ was deliberately set at the lower values in order to sputter in the stoichiometrically oxygen gas deficient conditions. Work function of the metal oxides was measured by cesium plasma immersion technique. The obtained min. work function values of AgO _x , NbO _x , WO _x , and TaO _x were 1.25, 1.38, 1.42, and 1.43 eV, resp. NbO _x and AgO _x are considered the most promising for a collector. A thermionic converter with a plane parallel type of a polycryst. W emitter and AgO _x collector, and an interelectrode spacing 0.1 mm at room temp., was set up and the power generation expts. were conducted. The max. power of 3.9 W/cm ² (voltage 0.6V; c.d. 6.5 A/cm ²) was obtained under the unignited mode operation at 1583 K. The barrier index was 1.5 V at 1578 K. Based on the exptl. results, a new type of a functionally graded material collector was proposed for a micro-gap thermionic converter.				
IT	Thermionic energy converters (development of refractory metal oxide collector materials and their thermionic converter performance)				
IT	Refractory metal oxides RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses) (development of refractory metal oxide collector materials and their thermionic converter performance)				
IT	1313-96-8, Niobium oxide	1314-35-8, Tungsten oxide, uses 1314-61-0, Tantalum oxide	20667-12-3, Silver oxide	RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses) (collector; development of refractory metal oxide collector materials and their thermionic converter performance)	
IT	7440-33-7, Tungsten, uses				

RL: DEV (Device component use); USES (Uses)
 (thermionic energy converter with **emitter** of; development of
 refractory metal oxide collector materials and their thermionic
 converter performance)

L3 ANSWER 16 OF 36 HCPLUS COPYRIGHT 2002 ACS

AN 1996:248255 HCPLUS

DN 124:304838

TI Manufacture of field-emission cold cathode

IN Yoshihara, Takuya

PA Nippon Electric Co, Japan

SO Jpn. Kokai Tokkyo Koho, 4 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI	JP 08031308	A2	19960202	JP 1994-182802	19940712
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AB The manuf. involves the following steps: (1) etching a deposited layer on a substrate to form the **emitter**, (2) coating the **emitter** with an insulating film and a gate electrode, (3) etching back to open the gate electrode, and (4) exposing the top of the **emitter**. The first deposited layer may be Ta. The insulating film may be an anodized film.

IT Cathodes

(field-emission, manuf. of field-emission cold cathode by etching **emitter**)

IT 7440-25-7, Tantalum, processes

RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses)

(**emitter**; manuf. of field-emission cold cathode by etching **emitter**)

IT 1314-61-0P, Tantalum oxide

RL: DEV (Device component use); PNU (Preparation, unclassified); PREP (Preparation); USES (Uses)

(insulating film; manuf. of field-emission cold cathode by etching **emitter**)

L3 ANSWER 17 OF 36 HCPLUS COPYRIGHT 2002 ACS

AN 1995:921960 HCPLUS

DN 123:327726

TI Capacitors, electrodes, or wiring structures for LSI

IN Numata, Ken; Aoki, Katsuhiro; Fukuda, Yukio; Nishimura, Akitoshi

PA Texas Instruments Inc., USA

SO Eur. Pat. Appl., 24 pp.

CODEN: EPXXDW

DT Patent

LA English

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI	EP 671768	A2	19950913	EP 1995-102002	19950214
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EP 671768	A3	19970820
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R: DE, FR, GB, IT, NL

JP 07226444	A2	19950822	JP 1994-39093	19940214
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US 5654567	A	19970805	US 1996-724159	19961001
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PRAI JP 1994-39093 19940214

US 1995-388330 19950214

AB In the capacitor, electrode, or wiring structure having an .alpha.-emitter (in particular, a Pt electrode), a shielding layer is provided, made of Ni, Co, Cu, and/or W, their compds. or alloys made of .gtoreq.2 of them, or compds. and alloys made of these metals and Si. It is possible to shield the .alpha.-particles effectively, to suppress generation of soft errors, and to enable use of Pt and other new materials in the electrodes and wiring, and to reduce the cost of the molding resin.

IT Electric capacitors
 Electrodes
 (for LSI)

IT Electric circuits
 (integrated, large-scale; .alpha.-particle shielding in capacitors and electrodes and wiring structures for)

IT Electric conductors
 (interconnections, for LSI)

IT 12587-46-1, Alpha particle
 RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses)
 (shielding against; in capacitors and electrodes and wiring structures for LSI)

IT 1314-61-0, Tantalum oxide 7440-02-0, Nickel, processes
 7440-06-4, Platinum, processes 7440-33-7, Tungsten, processes
 7440-48-4, Cobalt, processes 7440-50-8, Copper, processes 12060-59-2, Strontium titanate (SrTiO₃) 12626-81-2, Lead titanium zirconium oxide (PbTiO₃-ZrO₂-103) 37303-24-5, Barium strontium titanium oxide (Ba_{0.5} Sr_{0.5} TiO₃)
 RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses)
 (.alpha.-particle shielding in capacitors and electrodes and wiring structures contg.)

L3 ANSWER 18 OF 36 HCPLUS COPYRIGHT 2002 ACS
 AN 1995:895143 HCPLUS
 DN 123:318684
 TI High efficient radiation stable AlGaAs/GaAs solar cells with internal Bragg reflector
 AU Andreev, V. M.; Kalinovsky, V. S.; Komin, V. V.; Kochnev, I. V.; Lantratov, V. M.; Shvarts, M. Z.
 CS A. F. Phys.-Tech. Inst., St.-Petersburg, 194021, Russia
 SO Eur. Space Agency, [Spec. Publ.] ESA SP (1995), ESA SP-369(Vol. 2, Proceedings of the European Space Power Conference, 1995, Vol. 2), 367-70
 CODEN: ESPUD4; ISSN: 0379-6566
 DT Journal
 LA English
 AB The work presents an investigation of solar cells based on AlGaAs/GaAs heterostructures with internal Bragg reflector as back-surface reflectors grown by low-pressure MOCVD on n-GaAs substrates in horizontal resistively heated reactor. The typical structure consists of: Bragg reflector having 12 periods, n-GaAs base layer with thickness of 1.5-2.5 .mu.m, 0.4-0.5 .mu.m thick p-GaAs emitter, 0.07 .mu.m p-AlGaAs passivating window and top p-GaAs contact layers. The Bragg reflector with reflectance max. centered at wavelength 860 nm consists of twelve pair of AlAs/GaAs layers. Resulting Bragg reflector thicknesses have been 0.072 .mu.m for AlAs and 0.059 .mu.m for GaAs. In the case a peak of reflectance spectrum is close to unity in area 830-900 nm. This multi-layer quasi-dielectric stack would selectively reflect weakly absorbed photons with energies near the GaAs band gap for a second pass through the photoactive region increasing the photocurrent. The employment of the Bragg reflector allows to increase the external quantum efficiency in the

long wavelength of spectrum, to fabricate simultaneously thinner n-GaAs base layer and to increase the radiation resistance at 1 MeV and 3.75 MeV electron irradn. up to dose 1×10^{16} e/cm² and 3×10^{15} e/cm² correspondingly. The use of the internal Bragg reflector and Ta205 as antireflecting coating and prismatic cover allowed us to obtain efficiency 23.4% (17.7 suns, AM0, 25.degree.).

IT Photoelectric devices, solar
(high efficient radiation stable AlGaAs/GaAs solar cells with internal Bragg reflector)

IT Electron beam
(irradn. by; high efficient radiation stable AlGaAs/GaAs solar cells with)

IT Optical reflectors
(Bragg, internal; high efficient radiation stable AlGaAs/GaAs solar cells with)

IT 1314-61-0, Tantalum pentoxide
RL: TEM (Technical or engineered material use); USES (Uses)
(antireflection coating; high efficient radiation stable AlGaAs/GaAs solar cells with)

IT 1303-00-0, Gallium arsenide, uses 37382-15-3, Aluminum gallium arsenide (Al,Ga)As
RL: DEV (Device component use); USES (Uses)
(high efficient radiation stable AlGaAs/GaAs solar cells with)

L3 ANSWER 19 OF 36 HCPLUS COPYRIGHT 2002 ACS

AN 1995:769818 HCPLUS

DN 123:158267

TI Electron emitters

IN Tantani, Yasushi; Oota, Norio

PA Dainippon Printing Co Ltd, Japan

SO Jpn. Kokai Tokkyo Koho, 14 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI	JP 07094084	A2	19950407	JP 1993-259045	19930922
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AB ITO, Ta, Ta205 films, and Al cold cathodes are successively formed on glass substrates, resists are deposited on the whole surface, light projection from the bottom side and development leave the resists only in the neighborhood of the cathodes, insulator films (e.g., SiO₂) and gate electrode films are formed on the whole surface, and the resists are lifted, leaving the gate electrode films around the cathodes.

IT Cathodes
(field-emission, for electron emission)

IT 1314-61-0, Tantalum oxide (Ta205) 7429-90-5, Aluminum, uses 7440-25-7, Tantalum, uses 7631-86-9, Silica, uses 50926-11-9, Indium tin oxide
RL: DEV (Device component use); USES (Uses)
(electron emitters with cold cathodes)

L3 ANSWER 20 OF 36 HCPLUS COPYRIGHT 2002 ACS

AN 1995:613159 HCPLUS

DN 123:24173

TI Field-emission cathodes

IN Ito, Shigeo

PA Futaba Denshi Kogyo Kk, Japan

SO Jpn. Kokai Tokkyo Koho, 7 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 07094075	A2	19950407	JP 1993-260389	19930927
AB	The spindle-type field-emission cathodes (having corn-shaped emitters on a cathode electrode through a resistance layer) have conducting insulators between the resistance layer and the emitters, which are insulated by short-circuit current between the gate and the emitters. The conducting insulators may be formed on each emitter. The conducting insulators comprising Al, Ag, MnO ₂ , or (p-n junction) Se may be insulated through electromigration. The resistance layer may comprise amorphous Si or Ta ₂ O ₅ .				
IT	Cathodes (field-emission, field-emission cathodes with conducting insulator on emitters)				
IT	1313-13-9, Manganese dioxide, uses 7429-90-5, Aluminum, uses 7440-22-4, Silver, uses 7782-49-2, Selenium, uses RL: DEV (Device component use); USES (Uses) (field-emission cathodes with conducting insulator on emitters)				
IT	1314-61-0, Tantalum oxide (Ta ₂ O ₅) 7440-21-3, Silicon, uses RL: DEV (Device component use); USES (Uses) (resistance layers; field-emission cathodes with conducting insulator on emitters)				

L3 ANSWER 21 OF 36 HCAPLUS COPYRIGHT 2002 ACS

AN 1995:330517 HCAPLUS

DN 122:94649

TI Manufacture of field emission cathodes

IN Ito, Shigeo; Watanabe, Teruo; Ochiai, Hisataka; Ootsu, Kazuyoshi; Taniguchi, Masateru

PA Futaba Denshi Kogyo Kk, Japan

SO Jpn. Kokai Tokkyo Koho, 5 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE	
PI	JP 06124649	A2	19940506	JP 1992-270580	19921008	
AB	JP 3180466	B2	20010625	Title field emission cathode is manufd. by forming cathode conductor layer, insulator layer and gate layer on an insulating substrate, opening holes through the insulator and the gate layers, anodizing the cathode conductors in the holes to form resistance layers in an electrolyte in which the cathode conductor layer is used as the anode and a passive electrode as the cathode, then forming cone-shaped Mo emitters on the resistance layers. This simple manuf. process provide uniform independent resistance layer for each emitter.		
IT	Anodization (manuf. of field emission cathodes)					
IT	Cathodes (field-emission, manuf. of)					
IT	7439-98-7, Molybdenum, uses 7440-25-7, Tantalum, uses 7631-86-9, Silica, uses RL: DEV (Device component use); USES (Uses)					

(manuf. of field emission cathodes contg.)
 IT 1314-61-0, Tantalum oxide (ta2o5)
 RL: DEV (Device component use); TEM (Technical or engineered material use); USES (Uses)
 (manuf. of field emission cathodes contg.)

L3 ANSWER 22 OF 36 HCAPLUS COPYRIGHT 2002 ACS
 AN 1995:242453 HCAPLUS
 DN 122:21952
 TI Cathode **emitters** attached with a functional film
 IN Fujii, Masaru; Takizawa, Tomonori; Maeda, Mikako
 PA Mitsubishi Electric Corp, Japan
 SO Jpn. Kokai Tokkyo Koho, 9 pp.
 CODEN: JKXXAF

DT Patent
 LA Japanese

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 06139964	A2	19940520	JP 1992-281418	19921020
JP 2847003	B2	19990113		

AB A functional film adhered on the face of cathode **emitter** bulbs is a transparent plastic substrate coated with a functional layer which has .gtoreq.2 functions comprising antistatic elec. conducting, contrast-enhancement color-filtering, reflection-preventing, and/or surface-roughening properties.

IT Optical filters
 (adhered for cathode **emitter** bulb face)

IT Electric conductors
 (for antistatic film for cathode **emitter** bulb face)

IT Optical reflection
 (for cathode **emitter** bulb face)

IT Luminescent screens
 (cathodo-, multifunctional film adhered on tube face for)

IT 1314-23-4, Zirconium oxide (ZrO₂), processes 1314-61-0, Tantalum oxide (Ta₂O₅) 1314-98-3, Zinc sulfide (ZnS), processes 7631-86-9, Silica, processes 7783-40-6, Magnesium fluoride (MgF₂) 13463-67-7, Titanium oxide (TiO₂), processes

RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses)
 (reflection-preventive film for cathode **emitter** bulb face)

L3 ANSWER 23 OF 36 HCAPLUS COPYRIGHT 2002 ACS
 AN 1993:593897 HCAPLUS
 DN 119:193897
 TI Field-emission cathodes
 IN Ito, Shigeo; Watanabe, Teruo; Taniguchi, Masateru
 PA Futaba Denshi Kogyo Kk, Japan
 SO Jpn. Kokai Tokkyo Koho, 4 pp.
 CODEN: JKXXAF

DT Patent
 LA Japanese

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 05094760	A2	19930416	JP 1991-276233	19910930
JP 2720662	B2	19980304		

AB The field-emission cathodes comprise an insulative substrate, a cathodic electrode formed on the substrate, a thin Ta₂O₅ layer, as a resistor,

formed on the cathodic electrode, and an **emitter** connected to the cathodic electrode via through the resistor layer. The Ta₂O₅ layer has a uniform thickness since it can be formed by the anodic oxidn. method. The field-emission cathodes are used as electron sources for various display devices, sensors, etc.

IT Sensors
(field-emission cathodes for)

IT Cathodes
(field-emission, tantalum oxide resistor layer in, for display devices and sensors)

IT 1314-61-0, Tantalum pentoxide
RL: TEM (Technical or engineered material use); USES (Uses)
(resistor layer, in field-emission cathodes, for display devices and sensors)

L3 ANSWER 24 OF 36 HCPLUS COPYRIGHT 2002 ACS

AN 1993:542972 HCPLUS

DN 119:142972

TI Structure of high-efficiency solar cell and especially crystalline silicon solar cell

IN Schmidt, Wilfried; Wahl, Gerhard

PA Deutsche, Aerospace A.-G., Germany

SO Ger. Offen., 7 pp.

CODEN: GWXXBX

DT Patent

LA German

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	DE 4217428	A1	19930617	DE 1992-4217428	19920527
PRAI	DE 1991-4140488		19911209		

AB The solar cell for 1- or 2-side illumination has an n-p structure with p-type Si substrate and n-type **emitter** on the light-incident side. The solar cell of an optimal efficiency has on the light-incident side deep, on the surface low-doped n+ or p+ layers, integrated with higher-doped n++ or p++ layers on their region of ohmic contact surfaces of metallic contacts. The prepn. stages of the solar cell include deposition of absorption layers and Si oxide masking layers, removal of masking layers of the window opening regions by laser irradn. or etching with mixts. contg HF, and diffusion treatments.

IT Photoelectric devices, solar
(silicon, structure and manuf. of high-efficiency cryst.)

IT 12033-89-5, Silicon nitride, properties 13463-67-7, Titanium oxide, uses 59763-75-6, Tantalum oxide

RL: PRP (Properties)

(photoelec. solar cells contg. absorption layers of, cryst. silicon, structure and manuf. of high-efficiency)

IT 7440-21-3, Silicon, miscellaneous

RL: MSC (Miscellaneous)

(photoelec. solar cells, cryst., structure and manuf. of high-efficiency)

L3 ANSWER 25 OF 36 HCPLUS COPYRIGHT 2002 ACS

AN 1993:181663 HCPLUS

DN 118:181663

TI Field electron **emitters**

IN Komatsu, Hiroshi

PA Seiko Epson Corp., Japan

SO Jpn. Kokai Tokkyo Koho, 6 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 04280030	A2	19921006	JP 1991-40372	19910307
AB	A field electron emitter , which contains a protruded cathode for emitting electron, a gate electrode to apply an elec. field on the cathode, and an anode to collect emitted electron has a varistor which is series-connected with the cathode. The app. has stable electron flux.				
IT	Electron sources (emitters , field-based, contg. varistors)				
IT	Electric resistors (varistors, field electron emitters contg.)				
IT	59763-75-6, Tantalum oxide RL: DEV (Device component use); USES (Uses) (varistors from, field electron emitters contg.)				

L3 ANSWER 26 OF 36 HCAPLUS COPYRIGHT 2002 ACS

AN 1993:25051 HCAPLUS

DN 118:25051

TI Semiconductor base unit for solar cells

IN Eyckmans, Marc

PA Telefunken Systemtechnik GmbH, Germany

SO Ger. Offen., 4 pp.

CODEN: GWXXBX

DT Patent

LA German

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	DE 4110256	A1	19921001	DE 1991-4110256	19910328
AB	The unit of Si having a diffused high-ohmic emitter has a 3-layer cover with a plurality laser-inscribed slits. The cover layers consist of a 5-30-nm SiO ₂ or SiC base layer for passivating the semiconductor surface, a .apprx.70-nm TiO _x , Si ₃ N ₄ , or Ta ₂ O ₅ antireflection middle layer, and a SiO ₂ top layer. A 2nd low-ohmic emitter is diffused only in the slit regions.				
IT	Photoelectric devices, solar (silicon, with multilayered cover and laser-inscribed slits)				
IT	409-21-2, Silicon carbide, uses 1314-61-0, Tantalum pentoxide 7631-86-9, Silica, uses 12033-89-5, Silicon nitride, uses 13463-67-7, Titanium oxide (TiO ₂), uses RL: USES (Uses) (cover contg. layer of, silicon solar cells with multilayer)				
IT	7440-21-3, Silicon, uses RL: USES (Uses) (photoelec. solar cells, with multilayered cover and laser-inscribed slits)				

L3 ANSWER 27 OF 36 HCAPLUS COPYRIGHT 2002 ACS

AN 1990:244364 HCAPLUS

DN 112:244364

TI Semiconductor device having gold top layer-containing laminated electrode

IN Takabe, Akio

PA NEC Corp., Japan

SO Jpn. Kokai Tokkyo Koho, 2 pp.

CODEN: JKXXAF

antireflection coating, for solar cells)

IT Etching
(anisotropic, V-groove formation on gallium arsenide solar cells by)

IT Photoelectric devices, solar
(microgrooved, gallium arsenide, homojunction, with V-grooved front surface, performance of)

IT 1314-61-0, Tantalum pentoxide
RL: USES (Uses)
(antireflection coating, V-grooved gallium arsenide solar cell with)

IT 1303-00-0, Gallium arsenide, uses and miscellaneous
RL: USES (Uses)
(photoelec. solar cells, homojunction, with V-grooved front surface, performance of)

L3 ANSWER 29 OF 36 HCAPLUS COPYRIGHT 2002 ACS
AN 1990:88116 HCAPLUS
DN 112:88116
TI Stacked insulator structure thin-film electroluminescent display devices
AU Ohwaki, Junichi; Kozawaguchi, Haruki; Tsujiyama, Bunjiro
CS Opto-Electron. Lab., NTT, Tokai, 319-11, Japan
SO J. Electrochem. Soc. (1990), 137(1), 340-2
CODEN: JESOAN; ISSN: 0013-4651
DT Journal
LA English
AB A large-capacity, 704 .times. 1024 dot, thin film electroluminescent (TFEL) display panel with a d. of 4.2 lines/mm is fabricated by employing a stacked insulator structure. Improvement of the breakdown phenomenon, from propagation to self-healing mode, by interposing a self-healing type insulator between the metal electrode and the second Ta205 insulator is discussed. Emission efficiency and luminance are 1.6 and 1.8 times higher, resp., than with conventional TFEL devices. This is accomplished by application of a low-resistivity SiO2 film adjacent to the ZnS:Mn emitter.
IT Electroluminescent devices
(film, stacked insulator structure)
IT 1314-98-3, Zinc sulfide, uses and miscellaneous
RL: USES (Uses)
(electroluminescent display device with manganese-doped layer of, stacked insulator structure for)
IT 7439-96-5, Manganese, uses and miscellaneous
RL: USES (Uses)
(electroluminescent display device with zinc sulfide layer contg., stacked insulator structure for)
IT 7429-90-5, Aluminum, uses and miscellaneous 50926-11-9, ITO
RL: USES (Uses)
(electroluminescent thin-film display device with stacked insulator structure and layer of)
IT 1314-61-0, Tantalum oxide (Ta205) 7631-86-9, Silicon dioxide, uses and miscellaneous
RL: USES (Uses)
(insulator stacked structure contg., in thin-film electroluminescent display device)

L3 ANSWER 30 OF 36 HCAPLUS COPYRIGHT 2002 ACS
AN 1988:561913 HCAPLUS
DN 109:161913
TI A superconductor-semiconductor junction three-terminal device and its manufacture
IN Michigami, Osamu; Tanabe, Keiichi; Asano, Hidefumi; Kato, Yujiro; Kubo,

Shugo

PA Nippon Telegraph and Telephone Public Corp., Japan

SO Jpn. Kokai Tokkyo Koho, 8 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 63032974	A2	19880212	JP 1986-175485	19860728
	JP 07040612	B4	19950501		

AB The title devices include a superconducting base layer, a metal (a superconductor or an ordinary conductor) or semiconductor **emitter** layer contacting the base layer, and a compd. semiconductor collector layer on the other side of the base layer from the **emitter** layer; the collector layer comprises InAs_xSb_{1-x} or SbGa_xIn_{1-x} solid soln.; the collector and the **emitter** layers, the superconducting base layers, a tunnel barrier layer, and an insulating substrate are formed from lattice-matched materials; and the superconductor is selected from 1 of NbN, NbN-TiN, NbN-VN, NbN-ZrN, NbH-HfN, NbN-NbC, NbN-MoC, NbN-WC, NbC-MoC, NbC-WC, TaC, TaC-MoC, or TaC-WC. Preferably, the tunnel barrier layer or the substrate is selected from 1 of BaF₂, MnO, Mn₂O₃, MnO₂, TaO, GeO₂, Pb₃O₄.

IT 1310-53-8, Germanium oxide (GeO₂), uses and miscellaneous 1313-13-9, Manganese oxide (MnO₂), uses and miscellaneous 1314-41-6, Lead oxide (Pb₃O₄) 1317-34-6, Manganese oxide (Mn₂O₃) 1344-43-0, Manganese oxide (MnO), uses and miscellaneous 7787-32-8, Barium fluoride (BaF₂) 12035-90-4, Tantalum oxide (TaO) 12070-06-3, Tantalum carbide (TaC) 12274-94-1, Niobium carbide nitride (Nb(C,N)) 24621-21-4, Niobium nitride (NbN) 110770-52-0, Tantalum tungsten carbide ((Ta,W)C) 116738-93-3, Molybdenum niobium carbide nitride ((Mo,Nb)(C,N)) 116738-94-4, Niobium vanadium nitride ((Nb,V)N) 116738-95-5, Niobium titanium nitride ((Nb,Ti)N) 116738-96-6, Tantalum tungsten carbide (Ta_{0.61}W_{0.39}C) 116738-97-7, Molybdenum niobium carbide nitride (Mo_{0.15}Nb_{0.85}C_{0.15}N_{0.85}) 116738-98-8, Niobium carbide nitride (NbC_{0.3}N_{0.7}) 116738-99-9, Antimony indium arsenide (Sb_{0.32}InAs_{0.68}) 116739-00-5, Antimony indium arsenide (Sb_{0.39}InAs_{0.61}) 116739-01-6, Antimony indium arsenide (Sb_{0.46}InAs_{0.54}) 116740-05-7 116740-06-8

L3 ANSWER 31 OF 36 HCAPLUS COPYRIGHT 2002 ACS
AN 1987:128864 HCAPLUS
DN 106:128864
TI High temperature lamp coatings
AU Rancourt, James D.; Martin, Robert L., Jr.
CS Opt. Coat. Lab., Inc., Santa Rosa, CA, 95407, USA
SO Proc. SPIE-Int. Soc. Opt. Eng. (1986), 678(Opt. Thin Films 2: New Dev.), 185-91
CODEN: PSISDG; ISSN: 0277-786X
DT Journal
LA English
AB The efficacy of high efficiency lamps can be improved by recycling the unused IR energy. This is accomplished by reflecting the IR portion of the emitted energy back onto the **emitter** with a hot mirror coating on the lamp envelope. The envelopes of these lamps operate at high temps. so that special coating designs are used which function well at elevated temps. The coatings can also be used with other lamp applications, such as flashlamps and color correcting or selective lamps.
IT Electric lamps
(coatings for high temp. operation of)
IT Optical materials
(films, lamp, for high temp. operations)
IT Films
(optical, lamp, for high temp. operations)
IT 60676-86-0, Fused silica
RL: USES (Uses)
(lamp coating using substrate of, for high temp. operation)
IT 1314-61-0
RL: PRP (Properties)
(lamp coatings from silica and, for high temp. operation)
IT 7631-86-9, Silica, uses and miscellaneous
RL: USES (Uses)
(lamp coatings from tantalum, for high temp. operation)

L3 ANSWER 32 OF 36 HCAPLUS COPYRIGHT 2002 ACS
AN 1985:550611 HCAPLUS
DN 103:150611
TI Glow **emitters**. Compact energy source for IR radiation
AU Reinemann, Ulrich; Holzinger, Walter
CS Bayern-Chem. G.m.b.H., Ottobrunn, D-8012, Fed. Rep. Ger.
SO Int. Jahrestag. - Fraunhofer-Inst. Treib- Explosivst. (1985), 16th(Pyrotech.: Basic Princ., Technol., Appl.), 64/1-64/19
CODEN: IFTEDV; ISSN: 0722-4087
DT Journal
LA German
AB An IR glow **emitter** system for a semiautomatic flight guidance system was developed for antitank missiles HOT. The glow **emitter** element consists of thin-wall Ta pipes filled with a gasless pyrotech. heating mixt. (Pb3O4 + Si) which are localized concentrically on the perimeter of the missiles' rear tail. The pipes are heated during flight by the burning redox system and, upon reaching .apprx.1430.degree., their exothermic oxidn. by atm. O2 is initiated.
IT Guided missiles
(antitank, IR glow **emitter** for flight guidance system of)
IT Redox reaction
(of lead oxide and silicon, in pyrotech. heating mixt. for IR glow **emitter**, for antitank missile)
IT Oxidation

(of tantalum, in IR glow **emitter**, for antitank missile)

IT 1314-61-0P
 RL: FORM (Formation, nonpreparative); PREP (Preparation)
 (formation of, by oxidn. of tantalum IR glow **emitter**, for
 antitank missile)

IT 7440-25-7, uses and miscellaneous
 RL: USES (Uses)
 (in IR glow source, for antitank missile)

IT 7440-21-3, uses and miscellaneous
 RL: USES (Uses)
 (pyrotech. heating mixt. contg. lead oxide and, for IR glow
emitter, for antitank missile)

IT 1314-41-6
 RL: PRP (Properties)
 (pyrotech. heating mixt. contg. silicon and, in IR glow **emitter**
 , for antitank missile)

IT 7782-44-7, reactions
 RL: RCT (Reactant)
 (reaction of, with tantalum, in IR glow **emitter**, for antitank
 missile)

L3 ANSWER 33 OF 36 HCPLUS COPYRIGHT 2002 ACS
 AN 1978:607816 HCPLUS
 DN 89:207816
 TI Thermionic **emitters** based on rare earth element oxides and their
 analogs
 AU Kul'varskaia, B. S.
 CS USSR
 SO Radiotekh. Elektron. (1978), 23(9), 1989-94
 CODEN: RAEIA4; ISSN: 0033-8494
 DT Journal
 LA Russian
 AB Rare earth oxides, including their binary systems with oxides of metals of
 different valences, are an arsenal of new active materials for the
 electron sources of vacuum tubes and related devices. Figures are shown
 for the temp. characteristics of the emission current of Y2O3, rare earth
 oxides, double oxides, and systems involving Ta2O5. The dependence of
 band bending on donor concn. is shown.
 IT Rare earth oxides
 RL: USES (Uses)
 (as thermionic **emitters**)
 IT Electron donors
 (in rare earth oxides, band bending in relation to concn. of)
 IT Energy level, band structure
 (of rare earth oxides, effect of donor concn. on band bending in)
 IT Cathodes
 (thermionic, rare earth oxides in)
 IT 1306-38-3, uses and miscellaneous 1306-38-3D, solid solns. with tantalum
 oxide 1308-87-8 1312-81-8 1312-81-8D, solid solns. with yttrium
 oxide 1314-20-1D, solid solns. with praseodymium oxide 1314-23-4D,
 solid solns. with yttrium oxide 1314-36-9, uses and miscellaneous
 1314-36-9D, solid solns. with oxides 1314-61-0
 1314-61-0D, solid solns. with cerium oxide 12032-20-1
 12036-32-7 12036-32-7D, solid solns. with thorium oxide 12036-41-8
 12055-23-1D, solid solns. with yttrium oxide 12064-62-9
 RL: USES (Uses)
 (as thermionic **emitter**)

L3 ANSWER 34 OF 36 HCPLUS COPYRIGHT 2002 ACS

AN 1975:50395 HCPLUS
 DN 82:50395
 TI Emission constant of metallic refractory materials
 AU Haufler, G.
 CS Inst. Kernenerg., Univ. Stuttgart, Stuttgart, Ger.
 SO Dtsch. Luft- Raumfahrt, Mitt. (1973), DLR-Mitt. 73-29, 11-23
 CODEN: DLRMAT
 DT Report
 LA German
 AB The work function (.PHI.*) and the Richardson emission const. (AR) were
 detd. for polycryst. sintered samples of various compns. of the systems
 Zr-O, Zr-B, Zr-C, Ta-O, Nb-C, Zr-Ta-C, and Zr-Nb-C. The gas pressure was
 <10⁻⁸ torr, and the residual gas was analyzed continuously. In the binary
 Zr systems, .PHI.* decreases with increasing nonmetal content in the
 .beta.-Zr phase up to the solv. limit, then increases in the 2-phase
 region, and drops rapidly in the ZrC, etc., phase. Thus, for Zr-C at
 1750, .PHI.* = 4.01, 3.39, 4.15, and 2.67 eV for 0, 0.3 (solv. limit), 38
 (ZrC lower limit), and 45 at. % C, resp. The total decrease in .PHI.* at
 the .beta.-Zr solv. limit is the same for the 3 systems. In the Nb-C
 system, with increasing C content, .PHI.* decreases in the Nb phase,
 increases in the Nb + .beta.-Nb₂C region, decreases in the .beta.-Nb₂C
 phase, increases in the .beta.-Nb₂C + NbC region, and decreases in the NbC
 phase. The variations are smaller than those of the binary Zr systems.
 For any homogeneous phase of the systems studied, log AR depends linearly
 on .PHI.*. For a given metal, the various nonmetals yield the same AR vs.
 .PHI.* relation, indicating that only the metal electrons contribute to
 the emission. Under the same conditions, AR of Group IVB compds. is
 .apprxeq. 10 times that of Group VB compds. For all the systems, low
 .PHI.* values occur at phase limits of high nonmetal content. Esp.,
 ZrC0.82 is a promising cathode material with .PHI.* = 2.67 eV.
 Oil-diffusion pumps should be avoided in contact with these cathodes,
 since .PHI.* is strongly changed by C contamination of the emitter
 surface.
 IT Electron emission
 (from refractory transition metal compds.)
 IT Work function
 (of transition metal refractory compds.)
 IT Niobium carbide (NbC), solid solns. with zirconium carbide
 Tantalum carbide, solid solns. with zirconium carbide
 Zirconium carbide, solid solns. with refractory carbides
 RL: PRP (Properties)
 (electron emission from)
 IT 1314-61-0 12069-94-2 12741-10-5 51680-56-9 53801-45-9
 RL: PRP (Properties)
 (electron emission from)
 IT 7440-03-1, properties 7440-25-7, properties 7440-67-7, properties
 RL: PRP (Properties)
 (electron emission from compns. in systems of nonmetals and)
 IT 7440-42-8, properties 7440-44-0, properties 7782-44-7, properties
 RL: PRP (Properties)
 (electron emission from compns. in systems of transition metals and)
 L3 ANSWER 35 OF 36 HCPLUS COPYRIGHT 2002 ACS
 AN 1973:447075 HCPLUS
 DN 79:47075
 TI Bipolar transistors
 PA SESCOSEM-Societe Europeenne de Semiconducteurs et de Microelectronique
 SO Brit., 4 pp.
 CODEN: BRXXAA

DT Patent
 LA English
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	GB 1310487	A	19730321	GB 1971-11165	19710423
	FR 2085484	A1	19711224	FR 1970-15108	19700424
	FR 2085484	A5	19711224		
PRAI	FR 1970-15108		19700424		

AB A method for manuf. of bipolar transistors involving only 2 masking steps is described. Following conventional diffusion doping of Si for the base, collector, and **emitter**, contacts are applied, after removal of the dielec. layer, by 1st depositing a thermally oxidizable metal layer, e.g. Ta, Hf, or Zr, followed to deposition of a patterned layer of a metal which is insensitive to the heat treatment, e.g. Al. By positioning the Al at areas where contacts are desired, the underlying metal is not oxidized. The exposed metal is oxidized to a protective layer. Illustration in the case of Ta and Al involves oxidn. at 500.degree..

IT Transistors
 (silicon bipolar, tantalum-aluminum contacts in)
 IT Coating materials
 (tantalum oxide, in silicon bipolar transistors)
 IT Electric contacts
 (tantalum-aluminum, in silicon bipolar transistors)
 IT 1314-61-0
 RL: TEM (Technical or engineered material use); USES (Uses)
 (coatings, in silicon bipolar transistors)
 IT 7429-90-5, uses and miscellaneous
 RL: TEM (Technical or engineered material use); USES (Uses)
 (elec. contact from tantalum and, in silicon bipolar transistors)
 IT 7440-25-7, uses and miscellaneous
 RL: TEM (Technical or engineered material use); USES (Uses)
 (elec. contacts from aluminum and, in silicon bipolar transistors)

L3 ANSWER 36 OF 36 HCPLUS COPYRIGHT 2002 ACS
 AN 1972:545431 HCPLUS
 DN 77:145431
 TI Metal-oxide-semiconductorfield-effect transistor
 IN Croset, Michel; Nouailles, Noel
 PA SESCOSEM-Societe Europeenne de Semiconducteurs et de Microelectronique
 SO Ger. Offen., 10 pp.
 CODEN: GWXXBX

DT Patent
 LA German
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	DE 2101620		19720727		
PRAI	FR 1970-1565		19700116		

AB A substrate of a given cond. type is provided with **emitter** and collector regions of opposite cond. type by diffusion, and a dielec. layer as a gate region between them. The gate region consists, at least partially, of an oxide of a readily oxidizable metal and the **emitter** and collector regions are covered by metal pads. In a preferred version, the dielec. layer is Ta₂O₅ produced by oxidn. of a vapor-deposited Ta film in an O-contg. atm. at 500.degree. and the metal on the **emitter** and collector regions is Al. Oxide formed on the latter during heating is removed by etching. The substrate is Si. In the devices described, the dielec. layer has a high dielec. const. and good

04/09/2002

Serial No.:09/846,127

resistance to cosmic radiation.

IT Transistors
(field-effect, silicon-tantalum oxide-aluminum film structure,
radiation-resistant)

IT 1314-61-0
RL: USES (Uses)
(transistors of silicon MOS film structure with dielec. layer of,
radiation-resistant)

IT 7429-90-5, uses and miscellaneous
RL: USES (Uses)
(transistors of silicon with tantalum oxide dielec. films and collector
and **emitter** regions of)

File 2:INSPEC 1969-2002/Apr W1
 (c) 2002 Institution of Electrical Engineers

Set	Items	Description
S1	3811	CI=(TI SS(S) O SS) (S)NE=2
S2	259	CI=(TA SS(S) O SS) (S)NE=2
S3	7	CI=(W SS(S) SI SS(S) NI SS) (S)NE=3
S4	2	CI=(TA SS(S) AL SS(S) O SS(S) N SS) (S)NE=4
S5	54	CI=(TA SS(S) AL SS(S) O SS) (S)NE=3
S6	381	CI=(AL SS(S) O SS(S) N SS) (S)NE=3
S7	168415	PT OR PLATINUM OR AU OR GOLD
S8	75378	MOLYBDENUM OR MO OR TANTALUM OR TA
S9	110175	IRIDIUM OR IR OR RUTHENIUM OR RU
S10	75987	CHROMIUM OR CR
S11	26465	EMITTER? ? OR ECL OR (COLLECTOR(2N)ELECTRODE)
S12	441	(CATHODE (2N)LAYER) OR (FUSED(2N) ELECTROLYTE)
S13	2083	(TUNNEL?) (3N) (FILM? ? OR LAYER? OR COAT????)
S14	64303	(METAL???? OR ALLOY? OR AMALGAM? OR INGOT? OR BULLION?) (5N-) (DIELECTRIC? OR OXIDE OR FILM? ?)
S15	1463	TITANIUM(2N)MONOXIDE OR TITANIUM(2N)OXIDE
S16	0	TUNGSTEN (2N)NITRIDE(2N) SULFIDE
S17	0	ALUMINUM TANTALUM NITRIDE OXIDE
S18	9	ALUMINUM (2N)TANTALUM(2N)NITRIDE(2N)OXIDE
S19	27	ALUMINUM (2N)TANTALUM(2N)OXIDE
S20	122	ALUMINUM(2N)NITRIDE(2N)OXIDE
S21	31	TANTALUM OXIDE
S22	657	TANTALUM(2N)OXIDE
S23	24976	EMITTER? ? OR (COLLECTOR? ? (2N)ELECTRODE? ?)
S24	18	(S1 OR S15) AND S23
S25	2	(S2 OR S22) AND S23
S26	0	(S5 OR S19) AND S23
S27	1	(S6 OR S20) AND S23
S28	1895	(S7:S10) AND S23
S29	0	S28 AND S12
S30	10	S28 AND ((CATHODE) (2N) (FILM? ? OR LAYER? OR COAT????))
S31	49	S28 AND S14
S32	0	S31 AND S13
S33	4	((S1:S6) OR (S15:S22)) AND S13
S34	4	S33 NOT (S24 OR S25 OR S30)
S35	2	(S1 OR S15) AND ECL
S36	1	S35 NOT (S24 OR S25 OR S30)
S37	2	((S1:S6) OR (S15:S22)) AND ECL

24/3,AB/1
DIALOG(R)File 2:INSPEC
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7230759 INSPEC Abstract Number: B2002-05-1350P-004
Title: Increased THz **emitter** efficiency by coherent superposition in
a high repetition rate resonator
Author(s): Janke, C.; Bolivar, P.H.; Kurz, H.; Kunzel, H.
Author Affiliation: Inst. fur Halbleitertech. II, Rheinisch-Westfälische
Tech. Hochschule, Aachen, Germany
Conference Title: Technical Digest. Summaries of papers presented at the
Quantum Electronics and Laser Science Conference. Postconference Technical
Digest (IEEE Cat. No.01CH37172) p.201
Publisher: Opt. Soc. America, Washington, DC, USA
Publication Date: 2001 Country of Publication: USA 283+26
postdeadline papers pp.
ISBN: 1 55752 663 X Material Identity Number: XX-2001-02286
Conference Title: Technical Digest. Summaries of papers presented at the
Quantum Electronics and Laser Science Conference. Conference Edition
Conference Sponsor: APS/Div. Laser Sci.; IEEE Lasers & Electro-Opt. Soc.;
OSA-Opt. Soc. America
Conference Date: 6-11 May 2001 Conference Location: Baltimore, MD, USA
Language: English
Abstract: The wide range of attractive applications of time domain THz
sensing is strongly hampered by the low conversion efficiency of optically
excited THz **emitters**. Conversion efficiencies for most coherent THz
emitters are extremely low, typically on the order of 10⁻⁵.
Considerable interest is therefore directed towards possible ways of
increasing the THz **emitter** efficiency. We follow the approach of
exposing the **emitter** to a coherent phase-matched THz background field
during the emission process in order to radiate more energy before
dephasing occurs, thereby amplifying the amount of coherently emitted THz
radiation (Martini et al, 1998). In this presentation, we discuss recent
experiments where an InGaAs surface field **emitter** is placed in a high
repetition rate THz resonator. By synchronously pumping the **emitter**
with a high repetition rate (1 GHz) Ti:sapphire laser source (Bartels et
al, 1999), a net efficiency increase is demonstrated for the first time.
Subfile: B
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24/3,AB/2
DIALOG(R)File 2:INSPEC
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7230168 INSPEC Abstract Number: A2002-10-0762-004
Title: Design and performance of a THz emission and detection setup based
on a semi-insulating GaAs **emitter**
Author(s): Zhao, G.; Schouten, R.N.; van der Valk, N.; Wenckebach, W.Th.;
Planken, P.C.M.
Author Affiliation: Dept. of Appl. Phys., Delft Univ. of Technol.,
Netherlands
Journal: Review of Scientific Instruments vol.73, no.4 p.1715-19
Publisher: AIP,
Publication Date: April 2002 Country of Publication: USA
CODEN: RSINAK ISSN: 0034-6748
SICI: 0034-6748(200204)73:4L.1715:DPED;1-R
Material Identity Number: R017-2002-004

U.S. Copyright Clearance Center Code: 0034-6748/2002/73(4)/1715(5) /\$19.00

Language: English

Abstract: We have built a relatively simple, highly efficient, THz emission and detection system centered around a 15 fs Ti:sapphire laser. In the system, 200 mW of laser power is focused to a 120 μ m diam spot between two silverpaint electrodes on the surface of a semi-insulating GaAs crystal, kept at a temperature near 300 K, biased with a 50 kHz, +or-400 V square wave. Using rapid delay scanning and lock-in detection at 50 kHz, we obtain probe laser quantum-noise limited signals using a standard electro-optic detection scheme with a 1-mm-thick (110) oriented ZnTe crystal or a (110) oriented 0.1-mm-thick GaP crystal. The maximum THz-induced differential signal that we observe is $\Delta I/I = 7 \times 10^{-3}$, corresponding to a THz peak amplitude of 95 V/cm. The THz average power was measured to be about 40 μ W, to our knowledge, the highest power reported so far generated with Ti:sapphire oscillators as a pump source. The system uses off-the-shelf electronics and requires no microfabrication techniques.

Subfile: A

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24/3,AB/3

DIALOG(R)File 2:INSPEC

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7204060 INSPEC Abstract Number: A2002-08-4255R-016, B2002-04-4320G-069

Title: Ti:sapphire planar waveguide coherent broadband **emitter**

Author(s): Bhutta, T.; Salathe, R.P.; Shepherd, D.P.; Eason, R.W.; Pollnau, M.

Author Affiliation: Centre of Optoelectron. Res., Southampton Univ., UK

Conference Title: Technical Digest. Summaries of papers presented at the Conference on Lasers and Electro-Optics. Postconference Technical Digest (IEEE Cat. No.01CH37170) p.581-2

Publisher: Opt. Soc. America, Washington, DC, USA

Publication Date: 2001 Country of Publication: USA 604+72 post deadline papers pp.

ISBN: 1 55752 662 1 Material Identity Number: XX-2001-01869

Conference Title: CLEO 2001. Technical Digest. Summaries of papers presented at the Conference on Lasers and Electro-Optics. Postconference Technical Digest

Conference Sponsor: IEEE/Lasers & Electro-Opt. Soc.; OSA-Opt. Soc. America; Quantum Electron. Division of the Eur. Phys. Soc.; Opt. Soc. Japanese Quantum Electron. Joint Group

Conference Date: 6-11 May 2001 Conference Location: Baltimore, MD, USA

Language: English

Abstract: Summary form only given. In recent years, broadband fiber interferometers have become very popular as basic instruments used in optical coherence tomography (OCT) for imaging applications in the biomedical field. A major challenge in the further development and applicability of OCT has been the improvement of both its spatial resolution and dynamic range. The longitudinal resolution is inversely proportional to the optical bandwidth of the light source. Broadband luminescence from transition-metal-ion doped materials, (e.g., Ti:sapphire) can significantly improve the longitudinal resolution compared to superluminescent diodes (~30 nm FWHM), but the low brightness of its luminescence is insufficient for achieving a useful dynamic range in OCT. Femtosecond lasers have, therefore, been used as large-bandwidth high-brightness light sources, and subcellular imaging has recently been demonstrated in this way. However, current femtosecond light sources do not necessarily meet the requirements of compactness, ease of use, and low

cost. We present a simple broadband light source based on a Ti:sapphire planar waveguide. It operates in a wavelength region near 800 nm, applicable to the investigation of biotissue and detectable with simple silicon diodes, with a bandwidth comparable to that of a femtosecond light source.

Subfile: A B
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24/3,AB/4
DIALOG(R)File 2:INSPEC
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7184681 INSPEC Abstract Number: A2002-07-8160C-001, B2002-03-2550E-080
Title: Enhancing the surface passivation of $\text{TiO}_{\text{sub}} 2/$ coated silicon wafers
Author(s): Richards, B.S.; Cotter, J.E.; Honsberg, C.B.
Author Affiliation: Centre for Photovoltaic Eng., New South Wales Univ., Sydney, NSW, Australia
Journal: Applied Physics Letters vol.80, no.7 p.1123-5
Publisher: AIP,
Publication Date: 18 Feb. 2002 Country of Publication: USA
CODEN: APPLAB ISSN: 0003-6951
SICI: 0003-6951(20020218)80:7L.1123:ESPT;1-9
Material Identity Number: A135-2002-007
U.S. Copyright Clearance Center Code: 0003-6951/2002/80(7)/1123(3)/\$19.00
Language: English
Abstract: In this letter, we demonstrate good surface passivation of lightly diffused n-type solar cell emitters using titanium dioxide ($\text{TiO}_{\text{sub}} 2/$) thin films treated with a furnace oxidation process. Transient-photoconductance decay, X-ray photoelectron spectroscopy, and scanning electron microscopy measurements indicate that the silicon dioxide layer formed at the $\text{TiO}_{\text{sub}} 2/$:Si interface provides excellent surface passivation. Emitter dark saturation current densities of $4.7 \times 10^{14} \text{ A/cm}^2$ are achieved by this method, demonstrating that $\text{TiO}_{\text{sub}} 2/$ films are compatible with high-efficiency solar cell structures.

Subfile: A B
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24/3,AB/5
DIALOG(R)File 2:INSPEC
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6865664 INSPEC Abstract Number: A2001-08-4280A-002, B2001-04-4320M-002
Title: Bulk InAs mirror as a THz-radiation intra-cavity emitter in a femtosecond mode-locked Ti:sapphire laser
Author(s): Zhenlin Liu; Ono, S.; Otake, H.; Sarukura, N.; Tze-An Liu; Huang, K.F.; Ci-Ling Pan
Author Affiliation: Inst. for Molec. Sci., Okazaki, Japan
Conference Title: OSA Trends in Optics and Photonics. Advanced Solid State Lasers. Vol.34. Proceedings p.612-15
Editor(s): Injeyan, H.; Keller, U.; Marshall, C.
Publisher: Opt. Soc. America, Washington, DC, USA
Publication Date: 2000 Country of Publication: USA xvi+666 pp.
ISBN: 1 55752 628 1 Material Identity Number: XX-2000-00324
Conference Title: Proceedings of Topical Meeting on Advanced Solid-State Laser (ASSL 2000)
Conference Sponsor: Opt. Soc. America; IEEE/Lasers & Electro-Opt. Soc.;

Eur. Phys. Soc

Conference Date: 13-16 Feb. 2000 Conference Location: Davos, Switzerland

Language: English

Abstract: THz radiation is generated from bulk InAs shallow-incidence-angle mirror inside the cavity of a mode-locked Ti:sapphire laser. The magnetic field is also applied for radiation enhancement.

Subfile: A B

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? T S24/3,AB/6-18

24/3,AB/6

DIALOG(R)File 2:INSPEC

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6763314 INSPEC Abstract Number: A2000-24-4260F-024, B2000-12-4330B-059

Title: High-average-power, high-repetition-rate, femtosecond Ti:sapphire lasers with intra-cavity and extra-cavity CW-amplification schemes

Author(s): Liu, Z.; Murakami, H.; Kozeki, T.; Ono, S.; Otake, H.; Sarukura, N.

Author Affiliation: Inst. for Molecular Sci., Okazaki, Japan

Conference Title: Nonlinear Optics: Materials, Fundamentals, and Applications. Technical Digest. Postconference Edition. TOPS Vol.46 (IEEE Cat. No.00CH37174) p.295-7

Publisher: Opt. Soc. America, Washington, DC, USA

Publication Date: 2000 Country of Publication: USA xx+422 pp.

ISBN: 1 55752 646 X Material Identity Number: XX-2000-02452

Conference Title: Nonlinear Optics: Materials, Fundamentals, and Applications. Technical Digest. TOPS Vol.46

Conference Sponsor: Opt. Soc. America; IEEE/Lasers & Electro-Opt. Soc

Conference Date: 6-10 Aug. 2000 Conference Location: Kaua'i-Lihue, HI, USA

Language: English

Abstract: We have demonstrated high-average-power, high-repetition-rate, femtosecond Ti:sapphire lasers with newly-invented intra-cavity or extra-cavity CW-amplification schemes. The maximum output powers reach 3.4 W and 5.8 W, respectively. Such high average power and high repetition-rate femtosecond optical pulses will open up new application possibilities including parametric frequency conversion, THz-emitter driver, and material processing. This cw-amplification scheme will be also widely applicable to other low-gain and laser-pumped gain media.

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DIALOG(R)File 2:INSPEC

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6584429 INSPEC Abstract Number: B2000-06-1350P-018

Title: Efficient terahertz radiation generation from a bulk InAs mirror as an intracavity terahertz radiation emitter

Author(s): Zhenlin Liu; Ono, S.; Otake, H.; Sarukura, N.; Tze-An Liu; Kai-Fung Huang; Ci-Ling Pan

Author Affiliation: Inst. for Molecular Sci., Okazaki, Japan

Journal: Japanese Journal of Applied Physics, Part 2 (Letters) vol.39, no.4B p.L366-7

Publisher: Publication Office, Japanese Journal Appl. Phys,

Publication Date: 15 April 2000 Country of Publication: Japan

CODEN: JAPLD8 ISSN: 0021-4922

SICI: 0021-4922(20000415)39:4BL.1366:ETRG;1-P

Material Identity Number: C580-2000-008

Language: English

Abstract: Terahertz (THz) radiation is generated efficiently from a bulk InAs mirror with a shallow incidence angle inside the cavity of a femtosecond, mode-locked Ti:sapphire laser self-started by a strained saturable Bragg reflector. A magnetic field is also applied to the InAs mirror to enhance THz radiation.

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DIALOG(R)File 2:INSPEC

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6500197 INSPEC Abstract Number: B2000-03-4330B-046

Title: High precision multi-pass synchronously delay triggering control system

Author(s): Guo Liang-Fu; Zhang Xhu-Kui; Wang Xiao-Dong; Tang Jun; Zeng Xiao-Ming; Huang Xiao-Jun

Author Affiliation: Lab. for Laser Fusion, CAEP, Minyang, China

Journal: High Power Laser and Particle Beams vol.11, no.6 p.715-19

Publisher: Nucl. Soc. China,

Publication Date: 15 Dec. 1999 Country of Publication: China

CODEN: QYLIEL ISSN: 1001-4322

SICI: 1001-4322(19991215)11:6L.715:HPMP;1-V

Material Identity Number: D415-2000-001

Language: Chinese

Abstract: Precision synchronization of a high brightness facility is obtained and a delay triggering control signal is produced with wide range (1 ns~999 mu s) high precision (1 ns) adjustment and multi-pass output (20 passes) through the use of high performance fast-ECL circuit, standard frequency counting method and high precision linear saw tooth wave voltage generator.

Subfile: B

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DIALOG(R)File 2:INSPEC

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6423978 INSPEC Abstract Number: A2000-02-4255R-003, B2000-01-4320G-026

Title: Tunable two-pulse narrow-band laser **emitter** for lidar systems

Author(s): Korolev, V.I.; Mesnyankin, E.P.

Author Affiliation: Sci.-Res. Inst. for Comprehensive Tests on Opto-Electron. Instrum. & Syst., Vavilov (S.I.) State Opt. Inst., St. Petersburg, Russia

Journal: Kvantovaya Elektronika, Moskva vol.28, no.3 p.232-6

Publisher: Turpion Ltd.; Kvantovaya Elektronika,

Publication Date: Sept. 1999 Country of Publication: Russia

CODEN: KVEKA3 ISSN: 0368-7147

SICI: 0368-7147(199909)28:3L.232;1-B

Material Identity Number: C314-1999-012

Translated in: Quantum Electronics vol.29, no.9 p.787-91

Publication Date: Sept. 1999 Country of Publication: UK

CODEN: QUELEZ ISSN: 1063-7818

SICI of Translation: 1063-7818(199909)29:9L.787:TPNB;1-K

Language: English

Abstract: One of the ways of constructing a two-pulse laser **emitter** with a narrow spectral line, tunable over a wide wavelength range and based on a Ti:sapphire active medium, was investigated. This medium was excited by pulsed second-harmonic radiation of a neodymium glass laser. The parameters of the laser components were determined. These make it possible to construct a two-pulse source of radiation with a line width of Δ λ = 0.01 Å in each pulse, tunable in the range 670-1010 nm, and providing means for varying the spectral spacing between the pulses. The amplification of tunable laser radiation was investigated in a single- and four-pass Ti:sapphire amplifier with an axially excited active element 20 mm long.

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DIALOG(R)File 2:INSPEC

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6042464 INSPEC Abstract Number: A9822-7660-002

Title: FT-NMR detection of ^{45}Sc , ^{49}Ti and ^{93}Nb in TiO_2 single crystal

Author(s): Sato, K.; Takeda, S.; Fukuda, S.; Minamisono, T.; Tanigaki, M.; Miyake, T.; Maruyama, Y.; Matsuta, K.; Fukuda, M.; Nojiri, Y.

Author Affiliation: Dept. of Phys., Osaka Univ., Toyonaka, Japan

Journal: Zeitschrift fur Naturforschung, Teil A (Physik, Physikalische Chemie, Kosmophysik) Conference Title: Z. Nat.forsch. A, Phys. Phys. Chem. Kosmophys. (Germany) vol.53A, no.6-7 p.549-51

Publisher: Verlag der Zeitschrift fur Naturforschung,

Publication Date: June-July 1998 Country of Publication: Germany

CODEN: ZNASEI ISSN: 0932-0784

SICI: 0932-0784(199806/07)53A:6/7L.549:D449;1-0

Material Identity Number: E952-98007

Conference Title: XIVth International Symposium on Nuclear Quadrupole Interactions

Conference Date: 20-25 July 1997 Conference Location: Pisa, Italy

Language: English

Abstract: In order to determine the electric quadrupole moment of the short-lived beta **-emitter** ^{41}Sc from the quadrupole coupling constant in TiO_2 , we measured the field gradient by detecting the Fourier-Transformed-NMR of stable isotope ^{45}Sc doped in TiO_2 . Also, in order to study the electronic structure of impurities systematically, EFGs were measured for ^{45}Sc , ^{49}Ti and ^{93}Nb in a TiO_2 single crystal.

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DIALOG(R)File 2:INSPEC

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6042425 INSPEC Abstract Number: A9822-7660G-003

Title: Electric quadrupole interactions of the short-lived beta **-emitter** ^{12}N in insulator crystals (^{12}N implanted in single crystal TiO_2)

Author(s): Minamisono, T.; Sato, K.; Akai, H.; Takeda, S.; Maruyama, Y.; Matsuta, K.; Fukuda, M.; Miyake, T.; Morishita, A.; Izumikawa, T.; Nojiri, Y.

Author Affiliation: Dept. of Phys., Osaka Univ., Toyonaka, Japan
 Journal: Zeitschrift fur Naturforschung, Teil A (Physik, Physikalische Chemie, Kosmophysik) Conference Title: Z. Nat.forsch. A, Phys. Phys. Chem. Kosmophys. (Germany) vol.53A, no.6-7 p.293-300

Publisher: Verlag der Zeitschrift fur Naturforschung, Publication Date: June-July 1998 Country of Publication: Germany

CODEN: ZNASEI ISSN: 0932-0784

SICI: 0932-0784(199806/07)53A:6/7L.293:EQIS;1-Y

Material Identity Number: E952-98007

Conference Title: XIVth International Symposium on Nuclear Quadrupole Interactions

Conference Date: 20-25 July 1997 Conference Location: Pisa, Italy

Language: English

Abstract: The electronic structure of nitrogen atoms as impurities in an ionic $TiO_{2/}$ crystal has been investigated by analyzing electric field gradients (EFGs) measured by use of short-lived beta -emitting ^{12}N implanted following nuclear reactions. Conventional beta -NMR and its modification, suitable for the detection of quadrupole effects in the NMR spectra, were used for the investigation of hyperfine interactions of ^{12}N located in substitutional sites of O atoms and interstitial sites in the crystal. In order to deduce absolute values of the EFGs from the obtained eqQ/h , the quadrupole moment of ^{12}N has been determined from the NMR detection of ^{12}N implanted in BN(hexagonal) crystal. Here the EFG at the N atom in BN was measured by detecting the FT-NMR of ^{14}N in the crystal. The EFGs in $TiO_{2/}$ are compared with the theoretical predictions based on the ab initio band-structure calculation in the framework of the KKR method.

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DIALOG(R)File 2:INSPEC

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5996572 INSPEC Abstract Number: A9818-4250-037, B9809-4340-128

Title: Intracavity self-induced transparency of a multilevel absorber

Author(s): Muller, M.; Kalosha, V.P.; Herrmann, J.

Author Affiliation: Max Born Inst. fur Nichtlineare Opt. und Kurzzeitspektroskopie, Berlin, Germany

Journal: Physical Review A (Atomic, Molecular, and Optical Physics) vol.58, no.2 p.1372-81

Publisher: APS through AIP,

Publication Date: Aug. 1998 Country of Publication: USA

CODEN: PLRAAN ISSN: 1050-2947

SICI: 1050-2947(199808)58:2L.1372:ISIT;1-G

Material Identity Number: N687-98008

U.S. Copyright Clearance Center Code: 1050-2947/98/58(2)/1372(10) /\$15.00

Language: English

Abstract: Intracavity self-induced transparency of a three-level absorber is studied in the scope of solid-state laser generation of an ultrabroadband electromagnetic pulse that drives the population of all absorber levels through complete Rabi flopping. We show that at sufficient pump rates a Ti:sapphire laser forces an intracavity GaAs single quantum-well absorber, which provides an inter-valence-band transition in the THz domain in addition to two direct optical interband transitions,

into the self-induced transparency regime and acts as an all-solid-state ultrabroadband pulse **emitter**. In dependence on the resonator bandwidth, the intracavity pulse energy and the absorber dipole moments we obtain a multilevel self-induced transparency pulse spectrum which extends from the THz domain up to the ultraviolet. The steady-state sub-10-fs pulse consists of only a few optical cycles with the high-frequency components at its leading edge and a single to subcyclic THz component at its trailing edge.

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DIALOG(R)File 2:INSPEC
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5717933 INSPEC Abstract Number: A9722-7860F-002, B9711-4220-021
Title: Film electroluminescent **emitters** on rough substrates
Author(s): Gurin, N.T.; Sabitov, O.Yu.; Brigadnov, I.Yu.
Author Affiliation: Moscow State Univ., Ul'yanovsk, Russia
Journal: Pis'ma v Zhurnal Tekhnicheskoi Fizika vol.23, no.15-16 p.

7-12
Publisher: AIP,
Publication Date: Aug. 1997 Country of Publication: Russia
CODEN: PZTFDD ISSN: 0320-0108
SICI: 0320-0108(199708)23:15/16L.7;1-B
Material Identity Number: B283-97009
Translated in: Technical Physics Letters vol.23, no.8 p.577-9
Publication Date: Aug. 1997 Country of Publication: USA
CODEN: TPLEED ISSN: 1063-7850
SICI of Translation: 1063-7850(199708)23:8L.577:FEER;1-T
U.S. Copyright Clearance Center Code: 1063-7850/97/080577-3\$10.00
Language: English

Abstract: The authors present investigations of metal/semiconductor/composite-liquid insulator/metal (MSCM) electroluminescent structures deposited on ordinary smooth and rough glass substrates. M is the first transmitting $\text{SnO}/\text{sub 2}$ electrode with a second clamped metal electrode with micrometer-regulated movement; S is a $\text{ZnS}:\text{Mn}$ electroluminescent layer, and C is a layer of composite liquid insulator consisting of a mixture of PFMS-4 silicon-organic liquid and a barium titanate $\text{BaTiO}/\text{sub 3}$ powder filler.

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DIALOG(R)File 2:INSPEC
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5596492 INSPEC Abstract Number: B9707-3240C-006
Title: Dielectric-base transistors with doped channel
Author(s): Hato, T.; Yoshida, A.; Yoshida, C.; Suzuki, H.; Yokoyama, N.
Author Affiliation: Fujitsu Labs. Ltd., Atsugi, Japan
Journal: Applied Physics Letters vol.70, no.21 p.2900-2
Publisher: AIP,
Publication Date: 26 May 1997 Country of Publication: USA
CODEN: APPLAB ISSN: 0003-6951
SICI: 0003-6951(19970526)70:21L.2900:DBTW;1-Y
Material Identity Number: A135-97023

U.S. Copyright Clearance Center Code: 0003-6951/97/70(21)/2900/3/\$10.00

Language: English

Abstract: The dielectric-base transistor (DBT) is expected to be coupled with various functional oxides such as high-temperature superconductors and ferroelectrics. We experimented with lowering the conduction band of the channel to reduce the operating voltage. $\text{LaTiO}_{\text{sub}} 3$ deposited on $\text{SrTiO}_{\text{sub}} 3$ supplies carriers in the $\text{SrTiO}_{\text{sub}} 3$ substrate by displacing Sr^{2+} and La^{3+} . With this technique, we fabricated a $\text{YBa}_{\text{sub}} 2\text{Cu}_{\text{sub}} 3\text{O}_{\text{sub}} 7-x/\text{In}_{\text{sub}} 2\text{O}_{\text{sub}} 3/\text{SrTiO}_{\text{sub}} 3/\text{LaTiO}_{\text{sub}} 3/\text{SrTiO}_{\text{sub}} 3$ transistor with a partially doped channel. The transistor operates at under 1 V while maintaining a voltage amplification factor of 2, which is one order smaller than the 15 V operating voltage of a transistor with an undoped channel. The base potential relative to the **emitter** conduction band has been reduced to 0.3 eV.

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DIALOG(R)File 2:INSPEC

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5587261 INSPEC Abstract Number: B9707-3240C-001

Title: Electrical properties of $\text{Al}_{\text{sub}} 2\text{O}_{\text{sub}} 3/\text{(Ba,Rb)BiO}_{\text{sub}} 3/\text{SrTiO}_{\text{sub}} 3/\text{(Nb)}$ three terminal device

Author(s): Toda, F.; Yamada, T.; Hashimoto, K.; Abe, H.

Author Affiliation: Res. Lab., Oki Electr. Ind. Co. Ltd., Tokyo, Japan

Journal: Japanese Journal of Applied Physics, Part 1 (Regular Papers, Short Notes & Review Papers) Conference Title: Jpn. J. Appl. Phys. 1, Regul. Pap. Short Notes Rev. Pap. (Japan) vol.36, no.3B p.1990-3

Publisher: Publication Office, Japanese Journal Appl. Phys,

Publication Date: March 1997 Country of Publication: Japan

CODEN: JAPNDE ISSN: 0021-4922

SICI: 0021-4922(199703)36:3BL.1990:EPAB;1-W

Material Identity Number: F221-97007

Conference Title: 1996 International Conference on Solid State Devices and Materials (SSDM'96)

Conference Date: 26-29 Aug. 1996 Conference Location: Yokohama, Japan

Language: English

Abstract: A three terminal device with the $\text{Al}_{\text{sub}} 2\text{O}_{\text{sub}} 3/\text{(Ba,Rb)BiO}_{\text{sub}} 3/\text{Nb-doped SrTiO}_{\text{sub}} 3$ structure was fabricated using a superconducting base layer. The stable interface between $\text{(Ba,Rb)BiO}_{\text{sub}} 3$ and the artificial oxide barrier was obtained using *in situ* $\text{Al}_{\text{sub}} 2\text{O}_{\text{sub}} 3$ deposition. The output characteristics were measured as functions of input current. A current gain greater than 2 was obtained in the common **emitter** configuration.

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DIALOG(R)File 2:INSPEC

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5324977 INSPEC Abstract Number: A9617-7920H-001, B9609-2320-001

Title: Energy spectra of electrons emitted from samples with an internal electric field

Author(s): Olesik, J.; Calusinski, B.; Olesik, Z.

Author Affiliation: Inst. of Phys., Pedagogical Univ., Czestochowa,

Poland

Journal: Proceedings of the SPIE - The International Society for Optical Engineering Conference Title: Proc. SPIE - Int. Soc. Opt. Eng. (USA) vol.2638 p.94-102

Publisher: SPIE-Int. Soc. Opt. Eng,

Publication Date: 1995 Country of Publication: USA

CODEN: PSISDG ISSN: 0277-786X

SICI: 0277-786X(1995)2638L.94:ESEE;1-Z

Material Identity Number: C574-95263

U.S. Copyright Clearance Center Code: 0 8194 2004 2/95/\$6.00

Conference Title: Optical Characterization Techniques for High-Performance Microelectronic Device Manufacturing II

Conference Sponsor: SPIE

Conference Date: 25-26 Oct. 1995 Conference Location: Austin, TX, USA

Language: English

Abstract: This work is concerned with electron emission induced by an electric field and photoemission assisted by the field. The applied samples were emitters in the shape of semiconducting films evaporated on both sides of a glass substrate of thickness 0.2 mm. One side was an emitting surface whereas the other was a field electrode. The field electrode was supplied by a negative polarizing voltage $U_{\text{sub pol}}$. The emitting materials were $In_{\text{sub 2}}O_{\text{sub 3}}:Sn$ and titanium films. As a result of applying $U_{\text{sub pol}}$ and illumination, electrons and photoelectrons are released and enter the electron multiplier. Amplitude spectra of pulses were recorded by a multichannel pulse voltage analyser. Energy analysis of electrons released from the samples was performed by the retarding field method. Amplitude spectra at a given $U_{\text{sub pol}}$ and changing retarding field for titanium and oxide layers were compared. It was found that electron energy can exceed even 50 eV. For both types of films the influence of the illumination on the electron emission induced by an electric field was also investigated.

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DIALOG(R)File 2:INSPEC

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5172457 INSPEC Abstract Number: B9603-3240C-010

Title: Improved emitter-base junction with $In_{\text{sub 2}}O_{\text{sub 3}}$ in dielectric-base transistor

Author(s): Hato, T.; Takauchi, H.; Yoshida, A.; Tamura, H.; Fujimaki, N.; Oshima, Y.; Yokoyama, N.

Author Affiliation: Deposit Process Dev. Div., Fujitsu Labs. Ltd., Atsugi, Japan

Journal: Japanese Journal of Applied Physics, Part 1 (Regular Papers & Short Notes) vol.34, no.12A p.6379-81

Publisher: Publication Office, Japanese Journal Appl. Phys,

Publication Date: Dec. 1995 Country of Publication: Japan

CODEN: JAPNDE ISSN: 0021-4922

SICI: 0021-4922(199512)34:12AL.6379:IEBJ;1-F

Material Identity Number: C579-96001

Language: English

Abstract: We studied $In_{\text{sub 2}}O_{\text{sub 3}}$ for use as an intermediate layer between the YBCO emitter electrode and the $SrTiO_{\text{sub 3}}$ base region of dielectric-base transistors (DBT). We fabricated and tested a YBCO(001)/ $In_{\text{sub 2}}O_{\text{sub 3}}/(100)/SrTiO_{\text{sub 3}}/(110)$ heterostructure by using a laser-ablation deposition technique with high-density targets. Our

YBCO/In₂O₃/n-SrTiO₃/ diode structure showed a current density that was one order of magnitude larger than with an MgO intermediate layer.

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DIALOG(R)File 2:INSPEC

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4734693 INSPEC Abstract Number: B9409-3240C-028

Title: Increasing the current density of dielectric-base transistors with an MgO **emitter**-base barrier

Author(s): Yoshida, A.; Tamura, H.; Takauchi, H.; Hato, T.; Yokoyama, N.

Author Affiliation: Fujitsu Labs. Ltd., Atsugi, Japan

Journal: IEEE Transactions on Applied Superconductivity vol.4, no.2

p.76-80

Publication Date: June 1994 Country of Publication: USA

CODEN: ITASE9 ISSN: 1051-8223

U.S. Copyright Clearance Center Code: 1051-8223/94/\$04.00

Language: English

Abstract: We measured the current-voltage characteristics of YBa₂/Cu₃O_{7-x}/oxide/n-SrTiO₃/ diodes using NdGaO₃, LaAlO₃, CeO₂, and MgO as the oxide. MgO films had the highest current density. We then fabricated dielectric-base transistors with a YBa₂/Cu₃O_{7-x}/(YBCO) **emitter**/collector on a SrTiO₃ dielectric base with an MgO barrier. The transistors had both voltage and current gains exceeding unity at 4.2 K. The **emitter** current density was about 4×10^3 A/cm² at a collector-**emitter** voltage of 10 V and base-**emitter** voltage 10 V; this is 2 to 3 orders of magnitude larger than that of transistors with NdGaO₃ **emitter**-base barrier. We obtained a transconductance of around 0.4 mS at a collector-**emitter** voltage of 10 V for a device with a 6- μ m-diameter **emitter**.

Subfile: B

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DIALOG(R)File 2:INSPEC
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01821552 INSPEC Abstract Number: A82027668, B82018560
Title: Advances in high output voltage silicon solar cells
Author(s): Arndt, R.A.; Meulenberg, A.; Allison, J.F.; Weizer, V.G.
Author Affiliation: COMSAT Labs., Clarksburg, MD, USA
Conference Title: Fifteenth IEEE Photovoltaic Specialists Conference -
1981 p.92-6
Publisher: IEEE, New York, NY, USA
Publication Date: 1981 Country of Publication: USA 1471 pp.
Conference Date: 12-15 May 1981 Conference Location: Kissimmee, FL,
USA
Language: English
Abstract: Solar cells have been fabricated from 0.1 Omega -cm, p-type
silicon by a two-step diffusion process of **emitter** formation. The
cells are 200 mu m thick and 2*2 cm in area with a planar front surface
that has an anti-reflection (AR) coating of **tantalum oxide**.
Cr-Au-Ag contact metallization is on both sides of the cell. On the back,
the Cr-Au-Ag is applied over an aluminum diffused layer, and on the front,
it is applied through small holes in the AR coating. The best of these
cells has exhibited an open-circuit voltage of 654 mV under AM0
illumination.
Subfile: A B

30/3,AB/1
DIALOG(R)File 2:INSPEC
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6949961 INSPEC Abstract Number: A2001-14-7970-004, B2001-07-2320-011
Title: Electron field emission characterization of nanocrystalline diamond
thin film cold cathode devices

Author(s): Weiss, B.L.; Badzian, A.; Pilione, L.; Badzian, T.; Drawl, W.;
Morell, G.

Author Affiliation: Mater. Res. Lab., Pennsylvania State Univ.,
University Park, PA, USA

Conference Title: Amorphous and Nanostructured Carbon. Symposium
(Materials Research Society Symposium Proceedings Vol.593) p.227-31

Editor(s): Sullivan, J.P.; Robertson, J.; Zhou, O.; Allen, T.B.; Coll,
B.F.

Publisher: Mater. Res. Soc, Warrendale, PA, USA

Publication Date: 2000 Country of Publication: USA xvii+565 pp.

ISBN: 1 55899 501 3 Material Identity Number: XX-2000-02001

Conference Title: Amorphous and Nanostructured Carbon. Symposium

Conference Date: 29 Nov.-2 Dec. 1999 Conference Location: Boston, MA,
USA

Language: English

Abstract: Electron field emission measurements have been performed on
thin film cold cathode materials grown, on molybdenum, by
a modified MPACVD diamond process. Specifically the modification is due to
the addition of nitrogen and oxygen, in varying ratios, during the diamond
growth phase. Characterization using Raman spectroscopy shows features at
1190, 1330 and 1550 cm⁻¹. A simple triode device was fabricated for
electron emission characterization. KAPTON(R) film is used as the
insulating layer and a Mo mesh is used as the extraction gate
electrode. The collector is an indium tin oxide (ITO) coated
glass plate which is positively biased with respect to the gate electrode.
Field emission characteristics have shown current measurements of greater
than 1 microamp for fields of 40 V/micron. Gate currents are typically 1000
times greater than the emitted current. Issues currently being addressed
include improvement in the total emitted current, current stability and
device failure. We also present field emission measurements on diamond
films grown by HFCVD.

Subfile: A B

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6221394 INSPEC Abstract Number: A1999-10-7970-021, B1999-05-2320-033

Title: Planar field emitters fabricated by sulfur-doped boron
nitride

Author(s): Yokota, Y.; Tagawa, S.; Sugino, T.

Author Affiliation: Dept. of Electr. Eng., Osaka Univ., Japan

Journal: Journal of Vacuum Science & Technology B (Microelectronics and
Nanometer Structures) Conference Title: J. Vac. Sci. Technol. B,
Microelectron. Nanometer Struct. (USA) vol.17, no.2 p.642-6

Publisher: AIP for American Vacuum Soc,

Publication Date: March 1999 Country of Publication: USA

CODEN: JVTBD9 ISSN: 0734-211X

SICI: 0734-211X(199903)17:2L.642:PFEF;1-K

Material Identity Number: C067-1999-002
U.S. Copyright Clearance Center Code: 0734-211X/99/17(2)/642(5)/\$15.00
Conference Title: Fourth International Plasma-Based Ion Implantation
Workshop

Conference Date: 2-4 June 1998 Conference Location: Dearborn, MI, USA

Language: English

Abstract: Boron nitride (BN) films are grown on sapphire substrates by plasma-assisted chemical vapor deposition. BN films are doped with sulfur (S). The electrical resistivity of the S-doped BN film is reduced to 10^{13} Omega cm, while the electrical resistivity of the undoped BN film is 10^{11} Omega cm. It is demonstrated that the negative electron affinity appears on the BN surface. Insertion of a GaN layer between the BN film and sapphire leads to a tight adhesion of the BN film. Cathode and anode electrons are formed on the BN film and the sapphire substrate, respectively, by evaporating Ti and Au. An emission current of 1 mu A is obtained at an electric-field strength of 16 V/ mu m for the planar field emitter. An emission current density as high as 0.1 A/cm² is detected. It is expected that the planar field emitters can be operated at several tens V with a decreased cathode-anode spacing and that the present field emitter structure is applicable to a field-emission panel display.

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5960787 INSPEC Abstract Number: A9816-7970-001, B9808-2320-009
Title: Hafnium carbide films and film-coated field emission cathodes
Author(s): Mackie, W.A.; Tianbao Xie; Blackwood, J.E.; Williams, S.C.; Davis, P.R.
Author Affiliation: Linfield Res. Inst., McMinnville, OR, USA
Journal: Journal of Vacuum Science & Technology B (Microelectronics and Nanometer Structures) vol.16, no.3 p.1215-18
Publisher: AIP for American Vacuum Soc,
Publication Date: May-June 1998 Country of Publication: USA
CODEN: JVTBD9 ISSN: 0734-211X
SICI: 0734-211X(199805/06)16:3L.1215:HCFF;1-W
Material Identity Number: C067-98003
U.S. Copyright Clearance Center Code: 0734-211X/98/16(3)/1215(4)/\$15.00
Language: English

Abstract: We have previously reported on field emission improvements in turn-on voltages and emission stability using ZrC films as coatings on Si and Mo single emitters and emitter arrays. However, during our emission studies of bulk carbides, HfC was found to be slightly superior. We now report on work in progress investigating HfC films and HfC film coated field emission cathodes. Uses for arrays of these field emission cathodes range from video displays to microwave applications. This article deals with physical vapor deposition of HfC, absolute work function measurements, and electron emission properties of these film surfaces. This work demonstrates improvements by using HfC films over films of ZrC and an associated surface work function lowering of more than 1 eV in some instances compared to the clean surface.

Subfile: A B

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30/3,AB/4
 DIALOG(R)File 2:INSPEC
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5851482 INSPEC Abstract Number: A9808-7970-001, B9804-2320-008
 Title: Field electron emission from highly graphitic diamond films
 Author(s): Binglin Zhang; Ning Yao; Yunjun Li; Jintian He; Xiaoping Wang
 Author Affiliation: Dept. of Phys., Zhengzhou Univ., Henan, China
 Journal: Proceedings of the SPIE - The International Society for Optical
 Engineering Conference Title: Proc. SPIE - Int. Soc. Opt. Eng. (USA)
 vol.3184 p.225-8
 Publisher: SPIE-Int. Soc. Opt. Eng,
 Publication Date: 1997 Country of Publication: USA
 CODEN: PSISDG ISSN: 0277-786X
 SICI: 0277-786X(1997)3184L.225:FEEF;1-7
 Material Identity Number: C574-97226
 U.S. Copyright Clearance Center Code: 0277-786X/97/\$10.00
 Conference Title: Microelectronic Packaging and Laser Processing
 Conference Sponsor: SPIE; SPIE Singapore Chapter; Inst. Phys. Singapore;
 et al
 Conference Date: 25-26 June 1997 Conference Location: Singapore
 Language: English

Abstract: The diamond films were prepared by a microwave chemical vapor deposition system. **Molybdenum** substrates were used. The X-ray diffraction (XRD) spectra of the films contain peaks of the (111) and (220) facets of diamond. Scanning electron microscope (SEM) and optical micrograph reveal that the films consist of ball-like carbon structure, and diamond grains embedded on the balls. Raman spectra and surface resistance measurement also indicate that the films are highly graphitic diamond films. The field **emitter** was made of the diamond-ball like carbon **film cathode** and ITO anode. The transparent conducting anode technique was used to measure the I-V curves and the emission sites. The measurements were operated in a vacuum system with a base pressure of 10⁻⁴ torr. The turn-on field of 10 V/ μ m was obtained. After Ar⁺ ion impacting the highly graphitic diamond **film cathode**, the turn-on field was increased dramatically to 22 V/ μ m. A good quality diamond film **emitter** was also reported.

Subfile: A B
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30/3,AB/5
 DIALOG(R)File 2:INSPEC
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03868344 INSPEC Abstract Number: B91030912
 Title: Ir-coated dispenser **cathode** for CRT
 Author(s): Kimura, S.; Yakabe, T.; Matsumoto, S.; Miyazaki, D.; Yoshii, T.; Fujiwara, M.; Koshigoe, S.
 Author Affiliation: Toshiba Corp., Kawasaki, Japan
 Journal: IEEE Transactions on Electron Devices vol.37, no.12, pt.2
 p.2564-7
 Publication Date: Dec. 1990 Country of Publication: USA
 CODEN: IETDAI ISSN: 0018-9383
 U.S. Copyright Clearance Center Code: 0018-9383/90/1200-2564\$01.00
 Language: English
 Abstract: A compact dispenser cathode has been developed for application to cathode ray tubes (CRTs). A cathode **emitter**, comprising BaO, CaO, and Al₂O₃ in a molar ratio of 4:1:1, was impregnated into a

porous tungsten pellet. An intermetallic compound of tungsten and iridium was formed on the cathode pellet. Heater ratings were 6.3 V*0.2 A. Emission characteristics were measured by using color CRTs. As a result, a cathode peak loading of 15 A/cm² was ensured in the space-charge region. Furthermore, life tests with a peak loading of 7.5 A/cm² were conducted over 10000 h. The decrease in emission current after 10000 h was within only 10% of the initial value. Reliability of cathode performance was assured in terms of breakdown potential between the heater and the cathode, emission characteristics, life performance, grid emission, and the drift in cutoff potential. In addition, the effects of the coating thickness on the emission characteristics are discussed.

Subfile: B

30/3,AB/6
 DIALOG(R)File 2:INSPEC
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02768463 INSPEC Abstract Number: A86126838, B86067620
 Title: A study of surface damage in composite cathode materials in fast-flow CO₂ laser
 Author(s): Gnesin, G.G.; Levchenko, G.V.; Luban, R.B.; Barsuk, V.A.; Nesterenko, V.M.; Filimonov, M.Z.
 Author Affiliation: Inst. of Problems of Mater. Sci., Acad. of Sci., Kiev, Ukrainian SSR, USSR
 Journal: Poverkhnost'. Fizika, Khimiya, Mekhanika
 Publication Date: 1984 Country of Publication: USSR
 CODEN: PFKMDJ ISSN: 0207-3528
 Translated in: Physics, Chemistry and Mechanics of Surfaces vol.3, no.7 p.2110-24
 Publication Date: 1985 Country of Publication: UK
 CODEN: PCMSER ISSN: 0734-1520
 U.S. Copyright Clearance Center Code: 0734-1520/85/0307-2110\$20.00/0
 Language: English
 Abstract: Tests were carried out of a fast-flow electric-discharge CO₂ laser with a cathode unit containing electrode elements whose emitters are made of W/Cu- or Mo/Cu-based materials with LaB₆ inclusions. These materials were found to ensure a uniform discharge over the entire surface of the cathode unit. The cathode elements became coated with films formed by the emitter erosion and corrosion products. The density, thickness, and composition of these films varied depending on the composition of the emitter material. No short-circuiting of the electrode elements by the films occurred. The material containing 50% Mo, Cu, and inclusions of LaB₆ was the best and can be recommended for use as the emitter material in the cathode elements of fast-flow electric-discharge CO₂ lasers.

Subfile: A B

30/3,AB/7
 DIALOG(R)File 2:INSPEC
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01315046 INSPEC Abstract Number: A79019675, B79012541
 Title: A uniformly emitting MIM thin film cathode with evaporated insulator
 Author(s): Nisch, W.; Jonsson, C.
 Author Affiliation: Inst. fur Angewandte Phys., Univ. Tubingen, Tubingen,

West Germany

Journal: Optik vol.52, no.3 p.247-52

Publication Date: 1978 Country of Publication: West Germany

CODEN: OTIKAJ ISSN: 0030-4026

Language: German

Abstract: Describes the method of preparation of MIM (metal-insulator-metal) thin film cathodes of metal-Al₂O₃/Al₂O₃/Au type with evaporated insulator, and the study of the electrical properties and the electron emission from an emission electron microscope (EEM). The insulator was deposited by electron beam evaporation of ultrapure Al₂O₃ in clean vacuum (p=10⁻⁷ mbar). The insulator layers were 20 nm, and the cover electrodes 15 nm in thickness. The field-emission micrographs demonstrate the uniform emission over the emitter surface. Stable emission for hours was maintained with emission currents of 5*10⁻⁸ A over a surface of 8 mm² (current density about 6*10⁻⁷ A/cm²). Typically the total current between the base and the cover electrode was 2 mA at 8 V bias voltage.

Subfile: A B

30/3,AB/8

DIALOG(R)File 2:INSPEC

(c) 2002 Institution of Electrical Engineers. All rts. reserv.

00887752 INSPEC Abstract Number: A76030912

Title: Surface electron barriers for helium-3 and helium-4: experimental study of density dependence

Author(s): Broomall, J.R.; Onn, D.G.

Author Affiliation: Dept. of Phys. & Div. of Health Sci., Univ. of Delaware, Newark, DE, USA

Conference Title: Proceedings of the 14th International Conference on Low Temperature Physics Part I p.439-42

Editor(s): Krusius, M.; Vuorio, M.

Publisher: North-Holland, Amsterdam, Netherlands

Publication Date: 1975 Country of Publication: Netherlands xiii+528 pp.

ISBN: 0 7204 9302 1

Conference Date: 14-20 Aug. 1975 Conference Location: Otaniemi, Finland

Language: English

Abstract: In order to avoid the pressure limitations inherent in studying electrons at a liquid-vapor interface, hot electrons were injected directly into the helium fluid from the gold surface of a cold cathode (Al-oxide-Au) thin-film emitter. This device provides electrons of known energy distribution averaging about 1 eV. The injected current was collected at a gold-plated anode and measured as a function of atomic density of helium sample and applied electric field strength epsilon up to 60000 V/cm.

Subfile: A

30/3,AB/9

DIALOG(R)File 2:INSPEC

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00054915 INSPEC Abstract Number: B69017498

Title: Discharge tube satisfactorily low in radio-interfering noise

Inventor(s): Mizuno, M.; Akutsu, H.; Moriguchi, E.; Yamashita, K.; Kamiya, S.; Iwata, K.; Tawara, Y.; Iga, A.

Assignee(s): Matsushita Electronics Corporation, Osaka, Japan
Patent Number: US 3427492 Issue Date: 690211
Application Date: 670417
Priority Appl. Number: JP 41/25631 Priority Appl. Date: 660420
Country of Publication: USA
Language: English

Abstract: A discharge tube satisfactorily low in radio-interfering noise and equipped with cathodes coated with a cathode emitter consisting principally of oxides of barium, strontium and calcium and containing 0.05 to 10% by weight of a substance selected from the group consisting of cobalt boride and iron boride and further containing 1 to 8% by weight of a reducing metal having a high melting point and selected from the group consisting of zirconium, hafnium and tantalum, whereby a discharge tube low in noise intensity in the broadcasting frequency band is obtained.

Subfile: B

30/3, AB/10
DIALOG(R) File 2:INSPEC
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00054914 INSPEC Abstract Number: B69017497

Title: Discharge tube
Inventor(s): Mizuno, H.; Akutsu, H.; Moriguchi, E.; Yamashita, K.; Kamiya, S.; Iwata, K.; Tawara, Y.; Iga, A.
Assignee(s): Matsushita Electronics Corporation, Osaka, Japan
Patent Number: US 3427491 Issue Date: 690211
Application Date: 670417
Priority Appl. Number: JP 4125630 Priority Appl. Date: 660420
Country of Publication: USA
Language: English
Abstract: A discharge tube satisfactorily low in radio-interfering noise is obtained by using cathodes coated with a cathode emitter consisting principally of oxides of barium, strontium and calcium and containing iron-cobalt borides with part of the iron substituted by a substance selected from the group consisting of titanium, zirconium, hafnium, vanadium, niobium, tantalum, chromium, molybdenum, tungsten, aluminum and silicon and also containing a reducing metal having a high melting point selected from the group consisting of zirconium, hafnium, niobium and tantalum.

Subfile: B

FILE 'REGISTRY' ENTERED AT 14:45:16 ON 09 APR 2002

L1 1 S PT/CN
 L2 3 S AU/CN
 L3 2 S MO/CN
 L4 3 S TA/CN
 L5 0 S IR/CN
 L6 1 S IRIDIUM/CN
 L7 1 S RU/CN
 L8 3 S CR/CN

FILE 'HCAPLUS' ENTERED AT 14:46:16 ON 09 APR 2002

L9 440041 S PT OR PLATINUM OR AU OR GOLD
 L10 585655 S MOLYBDENUM OR MO OR TANTALUM OR TA
 L11 533406 S IRIDIUM OR IR OR RUTHERNIUM OR RU
 L12 459309 S CHROMIUM OR CR
 L13 32617 S Emitter OR ECL
 L14 4571 S (CATHODE (2N) LAYER) OR (FUSED(2N) ELECTROLYTE)
 L15 60 S ELETRON
 L16 4559 S (TUNNEL?) (3N) (FILM OR LAYER? OR COAT####)
 L17 225173 S (METAL#### OR ALLOY? OR AMALGAM? OR INGOT? OR BULLION?) (5N) (D
 L18 64246 S RUTHENIUM
 L19 5948 S L13 AND (L1-L12)
 L20 430 S L13 AND L18
 L21 38 S L19 AND L14
 L22 2 S L20 AND L14
 L23 0 S L22 NOT L21

FILE 'REGISTRY' ENTERED AT 14:52:07 ON 09 APR 2002

L24 718672 S AYS/CI

FILE 'HCAPLUS' ENTERED AT 14:52:25 ON 09 APR 2002

FILE 'REGISTRY' ENTERED AT 14:57:12 ON 09 APR 2002
 L25 14175 S (GOLD OR AU) AND L24

FILE 'HCAPLUS' ENTERED AT 14:58:10 ON 09 APR 2002
 L26 25335 S L25

FILE 'REGISTRY' ENTERED AT 15:00:25 ON 09 APR 2002
 L27 11186 S (PT OR PLATINUM) AND L24
 L28 23309 S L9 AND L24
 L29 168357 S L10 AND L24
 L30 8776 S L11 AND L24
 L31 3458 S RUTHENIUM AND L24
 L32 258173 S L12 AND L24

FILE 'HCAPLUS' ENTERED AT 15:03:52 ON 09 APR 2002
 L33 41 S L27 AND L13
 L34 154 S L28 AND L13
 L35 120 S L29 AND L13
 L36 22 S L30 AND L13
 L37 6 S L31 AND L13
 L38 200 S L32 AND L13
 L39 22 S L36 NOT L21
 L40 0 S L37 NOT L39
 L41 0 S L34 AND L14
 L42 1 S L34 AND L16

04/09/2002

Serial No.: 09/846,127

L43 410 S L34 OR L35 OR L38
L44 47 S L43 AND L17
L45 44 S L44 NOT (L39 OR L21)
L46 13 S L43 AND ((CATHODE?) (3N) (FILM OR LAYER? OR COAT####))
L47 0 S L46 NOT (L39 OR L21 OR L46)

L39 ANSWER 1 OF 22 HCAPLUS COPYRIGHT 2002 ACS
AN 2002:207357 HCAPLUS
DN 136:222998
TI Electron gun for technological linear accelerator
AU Khodak, I. V.; Kushnir, V. A.; Mitrochenko, V. V.; Perezhogin, S. A.; Stepin, D. L.; Zavada, L. M.; Zhiglo, V. F.
CS National Science Center, "ACCELERATOR" R and D Production Establishment, Kharkov Institute of Physics and Technology, Kharkov, Ukraine
SO Voprosy Atomnoi Nauki i Tekhniki, Seriya: Yaderno-Fizicheskie Issledovaniya (2000), (2), 86-88
CODEN: VANIEK
PB Natsional'nyi Nauchnyi Tsentr "Khar'kovskii Fiziko-Tekhnicheskii Institut"
DT Journal
LA English
AB The design of a diode electron gun for a powerful technol. electron linac and exptl. investigations of the beam parameters at the gun exit are considered. The gun features quick cathode replacement. The gun optics and beam parameters were calcd. using the EGUN code. The gun produces a beam current of 2 A at an anode voltage 25 kV. Measured beam parameters correspond to calcd. results.

L39 ANSWER 2 OF 22 HCAPLUS COPYRIGHT 2002 ACS
AN 2001:490559 HCAPLUS
DN 135:229286
TI Lifetime properties of the electrodes of a single-cell electricity generating channel in the Enisei thermionic nuclear power system
AU Koryukin, V. A.
CS Russian Science Center Kurchatov Institute, Russia
SO Atomic Energy (New York, NY, United States) (Translation of Atomnaya Energiya) (2001), 90(1), 1-11
CODEN: AENYEZ; ISSN: 1063-4258
PB Consultants Bureau
DT Journal
LA English
AB The working capacity and operating stability of multi- and single-cell electricity generating channels of the nuclear power systems TOPAZ-1 and -2 are largely detd. by the emission properties of the **emitter** and collector electrodes. The emission properties and elemental compn. of the electrode surfaces of single-cell EGC E-16MO, which exceeded their lifetimes in the exptl. systems Ya-24 and -81, which are prototypes of the Enisei space nuclear power system, sep. EGCs in the L channels, and in thermal stands, are detd. The previously performed comprehensive investigation of various processes on the electrode surfaces of EGC, and the database collected, made it possible to perform a statistical anal. of the results and to est. the lifetime stability of the electrodes on the basis of the emission properties up to 3 yr with 95% probability.

L39 ANSWER 3 OF 22 HCAPLUS COPYRIGHT 2002 ACS
AN 2001:396567 HCAPLUS
DN 134:375072
TI Efficient, cost-effective, long-lasting gas discharge lamp having an oxide **emitter** electrode
IN Gartner, Georg; Van den Hoek, Willem
PA Philips Corporate Intellectual Property G.m.b.H., Germany; Koninklijke Philips Electronics N.V.
SO Eur. Pat. Appl., 8 pp.
CODEN: EPXXDW
DT Patent

LA German

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	EP 1104005	A1	20010530	EP 2000-204015	20001114
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
	DE 19956322	A1	20010531	DE 1999-19956322	19991123
	JP 2001155679	A2	20010608	JP 2000-355169	20001122

PRAI DE 1999-19956322 A 19991123

AB A low-pressure discharge lamp consists of metal substrates and metal electrodes prep'd. from emitting metal powders made from reducing metals such as Al, Si, Ti, Zr, Hf, Ta, Mo, W and their alloys and noble metal powders selected from Re, Co, Ni, Ru, Pd, Rh, Ir, Pt and their alloys and .gt;eq.1 alk. earth oxide selected from CaO, SrO, and BaO.

IT 123-86-4, Butyl acetate 9004-70-0, Nitrocellulose

RL: NUU (Other use, unclassified); USES (Uses)
(binder; efficient, cost-effective, long-lasting gas discharge lamp having oxide **emitter** electrode)

IT 1304-28-5, Barium oxide, uses 1305-78-8, Calcium oxide, uses 1314-11-0, Strontium oxide, uses 1314-23-4, Zirconium dioxide, uses 7429-90-5, Aluminum, uses 7439-88-5, Iridium, uses 7439-98-7, Molybdenum, uses 7440-02-0, Nickel, uses 7440-05-3, Palladium, uses 7440-06-4, Platinum, uses 7440-15-5, Rhenium, uses 7440-16-6, Rhodium, uses 7440-18-8, Ruthenium, uses 7440-21-3, Silicon, uses 7440-25-7, Tantalum, uses 7440-32-6, Titanium, uses 7440-33-7, Tungsten, uses 7440-48-4, Cobalt, uses 7440-58-6, Hafnium, uses 7440-67-7, Zirconium, uses 42612-03-3

RL: DEV (Device component use); USES (Uses)
(efficient, cost-effective, long-lasting gas discharge lamp having oxide **emitter** electrode)

L39 ANSWER 4 OF 22 HCAPLUS COPYRIGHT 2002 ACS

AN 2001:16043 HCAPLUS

DN 134:186932

TI Change in the properties of the electrodes of thermionic single-element EGC at the initial stage of operation

AU Koryukin, V. A.

CS Russian Science Center Kurchatov Institute, Russia

SO Atomic Energy (New York) (Translation of Atomnaya Energiya) (2001), Volume Date 2000, 89(1), 555-564

CODEN: AENYEZ; ISSN: 1063-4258

PB Consultants Bureau

DT Journal

LA English

AB The initial emission-adsorption characteristics of the electrodes of single-element electricity-generating channel (EGC) thermionic energy converters were studied. The basic processes occurring at the electrodes and in the interelectrode space at the initial stage of operation, which ultimately affect the stability of the emission-adsorption characteristics of the electrodes, were examd. The dominant factors singled out are the evapn. of the **emitter** material and the transport of this material through the gap onto the collector and the effect of the active components of the residual gases (oxygen and carbon monoxide) in the interelectrode gap on the electrode properties. The surface state of the electrodes varies during the thermal vacuum prepn. and during placement of the EGC into the working regime as well as in the nominal regimes.

IT 7439-98-7, Molybdenum, processes 7440-44-0, Carbon, processes
 7440-46-2, Cesium, processes
 RL: ANT (Analyte); PEP (Physical, engineering or chemical process); ANST
 (Analytical study); PROC (Process)
 (adsorbate; change in properties of electrodes of thermionic
 single-element electricity-generating channel at initial stage of
 operation)

L39 ANSWER 5 OF 22 HCAPLUS COPYRIGHT 2002 ACS
 AN 2000:861257 HCAPLUS
 DN 134:36040
 TI Thermionic emission cathodes and their manufacture by impregnation
 IN Nonaka, Ikumitsu; Taguchi, Sadanori
 PA Hitachi, Ltd., Japan
 SO Jpn. Kokai Tokkyo Koho, 9 pp.
 CODEN: JKXXAF
 DT Patent
 LA Japanese
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 2000340097	A2	20001208	JP 1999-146639	19990526
AB	The cathode comprises porous sintered high m.p. metal substrates impregnated with electron emitting materials, with oxide layers formed in between the substrate and the impregnated materials. In manuf. of the cathodes by (a) impregnation of high m.p. porous metal substrate with an electron emitting material mainly consisting of alk. earth oxides, including Ba oxide, and alumina, (b) removal of the excess materials, and (c) heat treatment in vacuum; the substrate is oxidized prior to its impregnation with the material. Optionally a coating, having high m.p. and high work function (e.g. Os-Ru alloy) may be formed on the electron-emitting surface of the impregnated substrate. Cathodes, for cathode ray tubes, showing stable high c.d. and having long lifetime are obtained.				

IT 7440-33-7, Tungsten, processes 137949-21-4, Osmium 85, ruthenium
 15
 RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses)
 (manuf. of thermionic emission cathodes by surface oxidn. and
 impregnation of porous high m.p. metal sinters with aluminum barium
 mixed oxide electron emitters)
 IT 99035-55-9P, Barium calcium aluminate

L39 ANSWER 6 OF 22 HCAPLUS COPYRIGHT 2002 ACS
 AN 2000:823089 HCAPLUS
 DN 134:12467
 TI Indirect heating impregnated type cathode-ray tube
 IN Nonaka, Yasumitsu
 PA Hitachi, Ltd., Japan
 SO Jpn. Kokai Tokkyo Koho, 9 pp.
 CODEN: JKXXAF
 DT Patent
 LA Japanese
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 2000323010	A2	20001124	JP 1999-129119	19990510
AB	The cathode-ray tube has a high-m.p. metal cathode sleeve, a high-m.p.				

metal cup set on the top of the sleeve so that the sleeve is sealed with the cap, a heater inside the sleeve (for heating the cap), a metal funnel-shaped support, and a metal cylinder holder. The metal cup is filled with porous high-m.p. metal pellets impregnated with a hot-electron-emitting material based on Ba-contg. alk. earth metal oxide and the funnel-shaped support and the cylinder holder are placed so that thermal conduction from the heater through the sleeve and the cylinder holder is reduced, i.e., thermal efficiency is enhanced, and that the structure is not affected with vibration.

IT 99035-55-9, Aluminum barium calcium oxide
 RL: DEV (Device component use); USES (Uses)
 (electron **emitter**; in indirect heating impregnated type
 cathode-ray tube having funnel for supporting sleeve involving heater
 and cylinder holder)

IT 172890-99-2, Osmium, rutheni

L39 ANSWER 7 OF 22 HCAPLUS COPYRIGHT 2002 ACS

AN 2000:620816 HCAPLUS

DN 133:186817

TI Mo-(0-0.15)% Ir crystal for thermal **emitter** for surface ionization of organic compounds in air and method of its activation

IN Burkhanov, G. S.; Korenovskii, N. L.; Korolenko, I. I.; Kuz'mishchev, V. A.; Lyakishev, N. P.; Manokhin, I. K.; Nazarov, E. G.; Palitsin, V. V.; Prokhorov, A. M.; Rasulev, U. Kh.; Fesenko, A. V.; Chebyshev, A. V.; Shumilkin, A. V.

PA Institut Metallurgii im. A. A. Baikova RAN, Russia; Institut Elektroniki im. U. A. Arifova AN RUZ

SO Russ.

From: Izobreteniya 1999, (27), 393.

CODEN: RUXXE7

DT Patent

LA Russian

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI RU 2138877	C1	19990927	RU 1997-113927	19970812

AB Title only translated.

IT Electron sources

(molybdenum-iridium crystal for thermal **emitter** for surface ionization of org. compds. in air and method of activation)

IT Organic compounds, processes

RL: REM (Removal or disposal); PROC (Process)

(molybdenum-iridium crystal for thermal **emitter** for surface ionization of org. compds. in air and method of activation)

IT Ionization

(surface; molybdenum-iridium crystal for thermal **emitter** for surface ionization of org. compds. in air and method of activation)

IT 288309-86-4, Iridium 0-0.2, molybdenum 100

RL: TEM (Technical or engineered material use); USES (Uses)

(molybdenum-iridium crystal for thermal **emitter** for surface ionization of org. compds. in air and method of activation)

L39 ANSWER 8 OF 22 HCAPLUS COPYRIGHT 2002 ACS

AN 2000:241678 HCAPLUS

DN 132:259395

TI Cathode material of electron beam device and preparation method thereof

IN Choi, Jong Seo; Kim, Yoon Chang; Joo, Kyu Nam; Osaulenko, Nikolay; Shutovsky, Vladislav; Kultashev, Oleg

PA Samsung SDI Co., Ltd., S. Korea; Nikos-Eco, Ltd.

SO PCT Int. Appl., 23 pp.
CODEN: PIXXD2

DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2000021110	A1	20000413	WO 1999-KR599	19991001
	W: CN, DE, GB, JP, US				
	RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				
	KR 2000028717	A	20000525	KR 1999-41307	19990927
	EP 1129463	A1	20010905	EP 1999-970205	19991001
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, FI				
PRAI	UA 1998-105226	A	19981005		
	WO 1999-KR599	W	19991001		
AB	A cathode material of an electron beam device comprising 0.5 to 9.0% by wt. of a rare-earth metal of the Ce group, 0.5 to 15.0% by wt. of W and/or Re, 0.5 to 10% by wt. of Hf and the balance of Ir is provided. Since the cathode material has excellent plasticity, it is easy to manuf. small-size emitters. Also, since the d. of the electron emission of the cathode material is high and the working temp. is low, a long lifetime can be ensured. Also, the cathode material is useful as a cathode material of an electron beam device.				
IT	Alloying Cathodes Electronic device fabrication (cathode material of electron beam device and prepн. method thereof)				
IT	Rare earth alloys RL: DEV (Device component use); USES (Uses) (cerium-group; cathode material of electron beam device and prepн. method thereof)				

L39 ANSWER 9 OF 22 HCPLUS COPYRIGHT 2002 ACS
AN 2000:130857 HCPLUS
DN 132:139784
TI Electrical output parameters of efficient cylindrical thermionic energy converters
AU Kalandarishvili, A. G.; Kaibyshev, V. Z.; Lysikov, A. V.; Ermilov, B. I.
CS Kurchatov Institute, Russian Scientific Center, Russia
SO Atomic Energy (New York) (Translation of Atomnaya Energiya) (2000), Volume Date 1999, 87(2), 577-582
CODEN: AENYEZ; ISSN: 1063-4258
PB Consultants Bureau
DT Journal
LA English
AB Elec. output parameters are presented for four one-element cylindrical thermionic energy converters operating with cesium plasmas and electrode gaps from 0.3 to 0.8 mm. The cylindrical emitters were made by vapor-phase epitaxial deposition of tungsten on cylindrical molybdenum substrates having [111] axial orientation under the conditions where the deposit is automatically formed faced with (110) planes. The av. vacuum work function is up to 5.3 eV. The collector is made of a molybdenum-ruthenium alloy. Tests have been made on how the vacuum work function of the emitter influences the output parameters. Calcns. combined with expts. have been used to est. the work functions and electron temps. in the plasma.

04/09/2002

Serial No.: 09/846,127

IT Thermionic energy converters
(elec. output parameters of efficient cylindrical thermionic energy converters)
IT 69207-32-5
RL: DEV (Device component use); USES (Uses)
(collector; elec. output parameters of efficient cylindrical thermionic energy converters with)
IT 7440-33-7, Tungsten, uses
RL: DEV (Device component use); USES (Uses)
(emitter; elec. output parameters of efficient cylindrical thermionic energy converters with)
RE.CNT 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L39 ANSWER 11 OF 22 HCAPLUS COPYRIGHT 2002 ACS
AN 1997:105779 HCAPLUS
DN 126:232138
TI Ir-Ce cathodes as high-density **emitters** in electron-beam ion sources

STIC-EIC 2800 CP4-9C18

AU Rao, Roberto; Kultashev, Oleg
 CS Inst. Angewandte Phys. Frankfurt Main, Frankfurt, 60054, Germany
 SO Meas. Sci. Technol. (1997), 8(2), 184-188
 CODEN: MSTCEP; ISSN: 0957-0233
 PB Institute of Physics Publishing
 DT Journal
 LA English
 AB In electron-beam ion sources (EBIS/T), cathodes with high current densities are of great interest. Commonly used cathodes like impregnated tungsten cathodes and LaB6 cathodes emit current densities of 10 and 100 A cm-2, resp. In order to obtain high electron beam current densities of 103 A cm-2 for the stepwise ionization of few-electron, high-Z ions or for the fast prodn. of low-Z bare nuclei, electron-beam compression has to be very high using these cathodes. A high beam compression increases the beam temp., which limits the c.d. High current densities are also limited by the magnetic flux through the cathode. The application of small highly emissive cathodes, like Ir-Ce cathodes, reduces the magnetic flux through the cathode, so that a magnetic shield of the **emitter** is no longer required. By emitting relatively high currents from these small cathodes, hence emitted high emission c.d., the desired beam current densities can be attained by a moderate beam compression.

IT Ion sources
 (electron-beam; iridium-cerium cathodes as high-d. **emitters**
 in electron-beam ion sources)
 IT Cathodes
 (iridium-cerium cathodes as high-d. **emitters** in electron-beam
 ion sources)
 IT 39349-00-3
 RL: DEV (Device component use); USES (Uses)
 (iridium-cerium cathodes as high-d. **emitters** in electron-beam
 ion sources)

L39 ANSWER 12 OF 22 HCPLUS COPYRIGHT 2002 ACS
 AN 1997:94002 HCPLUS
 DN 126:112048
 TI Electron **emitter** and an electron source and image-forming
 apparatus using it
 IN Kishi, Fumio; Osada, Yoshiyuki; Kawade, Hisaaki; Tsukamoto, Takeo;
 Yoshida, Shigeki; Kusaka, Takao
 PA Canon K. K., Japan
 SO Eur. Pat. Appl., 45 pp.
 CODEN: EPXXDW
 DT Patent
 LA English
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	EP 747921	A2	19961211	EP 1996-303809	19960529
	EP 747921	A3	19961218		
	EP 747921	B1	19991215		
	R: DE, FR, GB, IT, NL				
	US 5939824	A	19990817	US 1996-654262	19960528
	CN 1146623	A	19970402	CN 1996-107769	19960530
	JP 09237568	A2	19970909	JP 1996-157431	19960530

PRAI JP 1995-154068 19950530
 JP 1995-342707 19951228

AB An electron **emitter** includes a pair of electrodes disposed
 opposite to each other, a conductive film in contact with both the pair of
 electrodes and an electron-emitting region formed in a part of the

conductive film. The conductive film is composed of fine particles including a 1st metal element serving as a main constituent element and a 2nd metal element. The 2nd metal element ppts. on the surface of the conductive film and thus forms a low-work-function layer. When a voltage is applied between the pair of electrodes, the 2nd metal element moves from the inside of the conductive film to at least part of its surface.

IT Cathodes
 (electron **emitter** and electron source and image-forming app.
 using it)
 IT Television
 (electron **emitter** and electron source for)
 IT Electron sources
 Electrooptical imaging devices
 (electron **emitter** for)
 IT 1314-08-5, Palladium oxide (PdO) 7440-02-0, Nickel, uses 7440-06-4, Platinum, uses 7440-32-6, Titanium, uses 7440-47-3, Chromium, uses 7440-57-5, Gold, uses 12006-69-8 12337-92-7 106524-62-3
 108673-66-1 122869-45-8, Palladium 95, titanium 5 (atomic) 127907-73-7
 185956-17-6 185956-20-1 185956-22-3 185956-25-6
 185956-28-9 185956-30-3 185956-32-5 185956-34-7 185956-36-9
 185956-38-1 185956-40-5 185956-42-7 185956-44-9 185956-46-1
 185956-48-3 185956-50-7 185956-52-9 185956-54-1 185956-56-3
 185956-58-5 185956-60-9 185956-62-1 185956-64-3 185956-66-5
 185956-69-8 185956-71-2 185956-73-4 185956-75-6 185956-77-8
 185956-79-0
 RL: DEV (Device component use); USES (Uses)
 (electron **emitter** contg.)

L39 ANSWER 13 OF 22 HCAPLUS COPYRIGHT 2002 ACS

AN 1996:483758 HCAPLUS

DN 125:127363

TI X-ray tubes with low-temperature **emitters**

IN Hell, Erich; Hoernig, Mathias; Kuhn, Helmut

PA Siemens A.-G., Germany

SO Ger., 8 pp.

CODEN: GWXXAW

DT Patent

LA German

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	DE 19513290	C1	19960725	DE 1995-19513290	19950407
	US 5703924	A	19971230	US 1996-627999	19960404
	CN 1138742	A	19961225	CN 1996-104604	19960405
	JP 08287854	A2	19961101	JP 1996-85536	19960408

PRAI DE 1995-19513290 19950407

AB X-ray tubes provided with an aperture positioned between the anode and the electron **emitter** so that the electrons must pass through it are described in which the aperture is maintained at the anode potential. Electrons produced at the anode are thus in a field-free region between the anode and the aperture, ensuring that only those electrons passing through the aperture reach the electron **emitter**. This reduces the no. of electron impacts on the **emitter** and thus extends the **emitter** (and thereby the tube) lifetime.

IT Group VIII element alloys

RL: DEV (Device component use); USES (Uses)

(x-ray tubes with apertures at anode potential for low-temp.
emitter preservation)

IT X-ray devices
 (sources, x-ray tubes with apertures at anode potential for low-temp.
 emitter preservation)

IT barium alloy, nonbase
 calcium alloy, nonbase
 cerium alloy, nonbase
 gadolinium alloy, nonbase
 lanthanum alloy, nonbase
 rhenium alloy, base
 thorium alloy, nonbase
 uranium alloy, nonbase
 yttrium alloy, nonbase
 RL: DEV (Device component use); USES (Uses)
 (x-ray tubes with apertures at anode potential for low-temp.
 emitter preservation)

IT 7439-98-7, Molybdenum, uses 7440-33-7, Tungsten, uses
 RL: DEV (Device component use); USES (Uses)
 (lanthanum oxide-doped; x-ray tubes with apertures at anode potential
 for low-temp. emitter preservation)

IT 12008-21-8, Lanthanum hexaboride 37186-87-1 39349-00-3
 39349-36-5, Iridium, lanthanum 39365-75-8, Thoriated tungsten
 RL: DEV (Device component use); USES (Uses)
 (x-ray tubes with apertures at anode potential for low-temp.
 emitter preservation)

IT 1312-81-8, Lanthanum oxide
 RL: DEV (Device component use); MOA (Modifier or additive use); USES
 (Uses)
 (x-ray tubes with apertures at anode potential for low-temp.
 emitter preservation)

L39 ANSWER 14 OF 22 HCAPLUS COPYRIGHT 2002 ACS
 AN 1996:142273 HCAPLUS
 DN 124:191717
 TI Small transformers and inverter and electron emitter circuits
 using thereof
 IN Yoshizawa, Katsuto; Abe, Tooru; Arakawa, Shunsuke
 PA Hitachi Metals Ltd, Japan
 SO Jpn. Kokai Tokkyo Koho, 8 pp.
 CODEN: JKXXAF
 DT Patent
 LA Japanese
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 07335450	A2	19951222	JP 1994-129046	19940610
AB	Title transformers comprise (1) a laminated magnetic core (thickness .ltoreq.3 mm) which is made from microcryst. (av. nanocryst. grain size .ltoreq.100 nm) or amorphous soft magnetic alloy films (thickness each .ltoreq.30 .mu.m), (2) a primary coil, and (3) secondary coil. The core is made of alloys contg. Fe and Co and/or Ni. The transformers using non-ferrite magnetic cores are compact and useful for inverter circuits in small cold cathode fluorescent lamps.				
IT	Transformers (fluorescent lamp; small transformers and inverter and electron emitter circuits using thereof)				
IT	Electric lamps (fluorescent, small; small transformers and inverter and electron emitter circuits using thereof)				
IT	Magnetic cores				

(transformer, laminated; small transformers and inverter and electron emitter circuits using thereof)

IT 174143-45-4 174143-47-6 174143-48-7 174143-49-8 174143-50-1
 174143-51-2 174143-52-3 174143-53-4 174143-54-5 174143-55-6
 174143-56-7 174143-57-8 174143-58-9 174143-59-0 174143-60-3
 174143-61-4 174143-62-5 174143-63-6 174143-64-7 174143-65-8
 174143-66-9 174143-67-0 174143-68-1 174143-69-2

RL: PRP (Properties)

(amorphous magnetic film for laminated core in small transformers and inverter and electron emitter circuits using thereof)

IT 174143-44-3 174143-46-5

RL: PRP (Properties)

(microcryst. magnetic film for laminated core in small transformers and inverter and electron emitter circuits using thereof)

L39 ANSWER 15 OF 22 HCPLUS COPYRIGHT 2002 ACS

AN 1995:350434 HCPLUS

DN 122:121061

TI Cathode substrates

IN Mizukami, Masahiko

PA Tokyo Tungsten Kk, Japan

SO Jpn. Kokai Tokkyo Koho, 3 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI JP 06176684 A2 19940624 JP 1992-349988 19921203

AB The substrates for electron emitters are made from sintered W particles whose surface is coated with Ir to give an Ir-W alloy. The alloy particles give decreased operational temp. and increased c.d.

IT Cathodes

(emitter; iridium-tungsten alloy substrate for)

IT 42612-03-3P

RL: DEV (Device component use); PNU (Preparation, unclassified); PREP (Preparation); USES (Uses)

(cathode emitter substrate; manuf. by sintering of powd. iridium and tungsten for)

IT 7440-33-7, Tungsten, reactions

RL: RCT (Reactant)

(powd.; sintering with iridium powder for formation of alloy cathode substrate)

IT 7439-88-5, Iridium, reactions

RL: RCT (Reactant)

(powd.; sintering with tungsten powder for formation of alloy cathode substrate)

L39 ANSWER 16 OF 22 HCPLUS COPYRIGHT 2002 ACS

AN 1994:287820 HCPLUS

DN 120:287820

TI Impregnated cathodes and manufacture thereof

IN Sugimura, Toshikazu; Narita, Kazunori

PA Kansai Nippon Electric, Japan

SO Jpn. Kokai Tokkyo Koho, 6 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 05282994	A2	19931029	JP 1992-77170	19920331
AB	The title process comprises dry pressing of a 1st powder mixt. from a refractory metal and an electron emitter , and a 2nd powder mixt. from the 1st powder mixt. and an additive of Ir, Os-Ru, Sc ₂ O ₃ , In ₂ O ₃ , and/or Y ₂ O ₃ to a lower and an upper portion of a form, resp., and sealing of the form into a capsule and hot isostatic pressing thereof.				
IT	Cathodes (sintered impregnated, manuf. of)				
IT	1304-28-5, Barium oxide, uses	1305-78-8, Calcium oxide, uses	1344-28-1, Aluminum oxide, uses		
	RL: PROC (Process) (impregnated sintered cathodes from emitters contg., manuf. of)				
IT	7440-33-7, Tungsten, uses				
	RL: PROC (Process) (impregnated sintered cathodes from, manuf. of)				
IT	1312-43-2P, Indium oxide (In ₂ O ₃)	1314-36-9P, Yttrium oxide (Y ₂ O ₃), uses	7439-88-5P, Iridium, uses	12060-08-1P, Scandium oxide (Sc ₂ O ₃)	87284-15-9P
	RL: IMF (Industrial manufacture); PREP (Preparation) (sintered cathodes impregnated with, manuf. of)				

L39 ANSWER 17 OF 22 HCAPLUS COPYRIGHT 2002 ACS

AN 1994:181090 HCAPLUS

DN 120:181090

TI Manufacture of impregnated cathode **emitter**
IN Suzuki, Yukio; Taguchi, Tadanori; Saito, Shunji
PA Hitachi Ltd, Japan
SO Jpn. Kokai Tokkyo Koho, 4 pp.
CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 05225897	A2	19930903	JP 1992-26514	19920213
AB	The title manufg. involves (1) heating a sintered porous refractory metal and an electron-emitting metal carbonate press-molded pellet in a reductive atm. to decomp. the carbonate until completion of CO ₂ discharging and also reducing oxide surface of the porous metal, (2) laminating the donor metal on the porous refractory metal in-situ in the reductive atm., and (3) heating the laminate at a high temp. to melt the press-mold to be impregnated into the porous refractory metal. The manufg. process provides the impregnation in-situ in the reductive atm. for preventing deterioration of emission characteristics.				
IT	Cathodes (thermionic, manuf. of, by impregnation of emitter material melts in porous metal supports)				
IT	87284-15-9, Osmium, ruthenium	RL: TEM (Technical or engineered material use); USES (Uses) (coatings, on cathode emitters)			
IT	471-34-1, Calcium carbonate, uses	1344-28-1, Alumina, uses	513-77-9, Barium carbonate		
	RL: USES (Uses) (electron emission material contg., for impregnation in porous refractory metal supports)				
IT	7440-33-7, Tungsten, uses				

RL: USES (Uses)
 (porous support for cathode **emitter**, impregnation of
emitter material in)

L39 ANSWER 18 OF 22 HCPLUS COPYRIGHT 2002 ACS
 AN 1993:676705 HCPLUS
 DN 119:276705
 TI Sub-surface iridium depletion in dilute solution tungsten-iridium alloys
 due to high temperature work function testing
 AU D'Cruz, L. A.; Jacobson, D. L.
 CS Dep. Chem., Bio Mater. Eng., Arizona State Univ., Tempe, AZ, 85287, USA
 SO Int. J. Refract. Met. Hard Mater. (1993), Volume Date 1992, 11(4), 223-34
 CODEN: IRMME3; ISSN: 0958-0611
 DT Journal
 LA English
 AB W-Ir alloys contg. <2 wt.% Ir exhibit the highest work functions among
 candidate alloys for thermionic **emitters**. The interrelationship
 between work function, compn., and microstructure of the alloys was
 investigated to ascertain the role of Ir on modification of the work
 function. The effective work function was detd. using a vacuum emission
 vehicle employing the thermionic method. Electron probe microanal. was
 used to characterize the sub-surface Ir depletion caused by elevated temp.
 testing. Theor. ests. of the equil. segregation of Ir in W could not
 adequately account for the magnitude of Ir depletion found at the
 electrode surfaces. An alternative explanation for the sub-surface Ir
 depletion is proposed and involves the creation of Ir-rich surface layers
 during the pre-test heat treatment of the alloys.
 IT Work function
 (of tungsten-iridium alloys, subsurface iridium depletion during
 high-temp. testing of)
 IT 133754-99-1 151530-83-5, Iridium 0.2, tungsten 100
 151530-84-6, Iridium 1.7, tungsten 98
 RL: PRP (Properties)
 (iridium loss from, during high-temp. work function testing)
 IT 7439-88-5, Iridium, properties
 RL: PRP (Properties)
 (loss of, from tungsten-iridium alloys, during high-temp. work function
 testing)

 L39 ANSWER 19 OF 22 HCPLUS COPYRIGHT 2002 ACS
 AN 1993:114902 HCPLUS
 DN 118:114902
 TI Impregnated cathode and cathode-ray tube using it
 IN Nonaka, Ikumitsu
 PA Hitachi, Ltd., Japan
 SO Jpn. Kokai Tokkyo Koho, 8 pp.
 CODEN: JKXXAF
 DT Patent
 LA Japanese
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 04280029	A2	19921006	JP 1990-138181	19900530

 AB The cathode is obtained by melting an electron-emitting alk. earth oxide
 contg. at least BaO and CaO at mol ratio 0.5-1.5 in a porous substrate of
 W, Mo, Ta, Re, Ir, etc., or their alloys and impregnating. The cathode
 showed good emission lifetime.
 IT Cathode-ray tubes
 (impregnated cathodes for, contg. alk. earth oxides in porous

refractory metals)
 IT Cathodes
 (impregnated, contg. alk. earth metal oxides, for cathode-ray tubes)
 IT 12060-08-1, Scandium oxide 1304-28-5, Barium oxide, uses 1305-78-8,
 Calcium oxide, uses
 RL: PROC (Process)
 (cathodes from refractory metals impregnated with, for cathode-ray
 tubes)
 IT 7439-88-5, Iridium, uses 7439-98-7, Molybdenum, uses 7440-04-2,
 Osmium, uses 7440-15-5, Rhenium, uses 7440-18-8, Ruthenium, uses
 7440-25-7, Tantalum, uses 7440-33-7, Tungsten, uses 87284-15-9
 RL: DEV (Device component use); USES (Uses)
 (cathodes from, impregnated with alk. earth oxides, for cathode-ray
 tubes)
 IT 1344-28-1, Alumina, uses
 RL: USES (Uses)
 (impregnated cathode electron **emitter**, for cathode-ray tube)

L39 ANSWER 20 OF 22 HCAPLUS COPYRIGHT 2002 ACS

AN 1991:693103 HCAPLUS

DN 115:293103

TI Impregnated cathodes for large-density current

IN Watabe, Isato; Yamamoto, Yoshihiko; Sasaki, Susumu; Yaguchi, Tomio

PA Hitachi, Ltd., Japan

SO Jpn. Kokai Tokkyo Koho, 4 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI	JP 03105827	A2	19910502	JP 1989-242006	19890920
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AB A long-life cathode, comprising a thermally resistive porous metal base impregnated with an electron-emitting substance, and a thin film from a mixt. of W, Sc₂O₃, and Sc₂W₃O₁₂ (or Sc₆W₀12), which decreases the work function of the electron-emission surface of the metal base, is characterized in that a Sc-supplying thin intermediate layer is placed between the metal base and the thin film.

IT Cathodes

(impregnated, for large-d. current)

IT 89742-05-2

RL: USES (Uses)

(electron-**emitter**, cathodes impregnated with, for large-d.
 current)

IT 7440-33-7, Tungsten, uses and miscellaneous 12060-08-1, Scandium oxide
 (Sc₂O₃) 12293-96-8, Scandium tungsten oxide (Sc₆W₀12) 13701-71-8,
 Scandium tungsten oxide (Sc₂W₃O₁₂) 137774-52-8 137774-53-9
 137774-54-0 137774-55-1 137774-56-2 137774-57-3
 137774-58-4 137774-59-5 137774-60-8 137774-61-9 137774-62-0

RL: USES (Uses)

(impregnated cathodes contg., for large-d. current)

L39 ANSWER 21 OF 22 HCPLUS COPYRIGHT 2002 ACS
 AN 1991:93176 HCPLUS
 DN 114:93176
 TI Iridium-coated dispenser cathode for CRT
 AU Kimura, Sakae; Yakabe, Toru; Matsumoto, Sadao; Miyazaki, Daisuke; Yoshii, Tsuyoshi; Fujiwara, Minoru; Koshigoe, Shimpei
 CS Electron Device Div., Toshiba Corp., Saiwaiku, 210, Japan
 SO IEEE Trans. Electron Devices (1990), 37(12, Pt. 2), 2564-7
 CODEN: IETDAI; ISSN: 0018-9383
 DT Journal
 LA English
 AB A compact dispenser cathode was developed for application to cathode ray tubes (CRT). A cathode **emitter** comprised BaO, CaO, and Al₂O₃ in a molar ratio of 4:1:1, was impregnated into a porous W pellet. Intermetallic compd. of W and Ir was formed on the cathode pellet. Heater ratings were 6.3 V x 0.2 A. Emission characteristics were measured by using color CRT's. As the result, a cathode peak loading of 15 A/cm² was ensured in the space-charge region. Life tests with a peak loading of 7.5 A/cm² were conducted over the 10000 h. The decrease in emission current after 10000 h was within only 10% of the initial value. Reliability of cathode performance was assured in terms of breakdown potential between the heater and the cathode, emission characteristics, life performance, grid emission, and the drift in cutoff potential. The effects of the coating thickness upon the emission characteristics are discussed.
 IT Cathodes
 (dispenser, iridium coated, for cathode ray tube)
 IT 7439-88-5, Iridium, uses and miscellaneous
 RL: USES (Uses)
 (cathodes coated with, for cathode ray tubes)
 IT 7440-25-7, Tantalum, uses and miscellaneous 37377-76-7
 39332-67-7, Kovar
 RL: USES (Uses)
 (cathodes contg., emission characteristics of)
 IT 89742-05-2, Barium calcium aluminate (Ba₄CaAl₂O₈)
 RL: USES (Uses)
 (cathodes contg., impregnated in tungsten pellet, emission characteristics of)
 IT 7440-33-7, Tungsten, uses and miscellaneous
 RL: USES (Uses)
 (cathodes of barium calcium aluminate impregnated in pellets of, emission characteristics of)

L39 ANSWER 22 OF 22 HCPLUS COPYRIGHT 2002 ACS
 AN 1986:198276 HCPLUS
 DN 104:198276
 TI Semiconductor device with a tantalum-iridium barrier layer contact structure
 IN Todd, Anthony G.; Wickenden, Dennis K.
 PA General Electric Co. PLC, UK
 SO U.S., 4 pp.
 CODEN: USXXAM
 DT Patent
 LA English
 FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI US 4546373	A	19851008	US 1984-575066	19840130
PRAI GB 1983-3298		19830207		

AB In a bipolar transistor for a temp. sensor having an adjustable const.-current source connected between the base and collector regions, and a high-gain amplifier having the input connected to the current source and the output to the **emitter** region, the **emitter**-base junction is a heterojunction, Au contacts are sep'd. by amorphous Ta-Ir alloy barrier layers from the n-type GaAs base, the p-type GaAs collector, and the p-type Ga_{1-x}Al_xAs **emitter**, resp. The compn. range of the alloy is 40-70% Ta, and the lower end of this range supplies a better barrier to Au migration. Hence, the sensor can be used to measure higher temps.

IT Temperature
(sensors, heterojunction bipolar transistors for, contg. iridium-tantalum alloy diffusion barriers)

IT Transistors
(bipolar, contg. heterojunctions and iridium-tantalum alloy diffusion barriers, for temp. sensors)

IT 1303-00-0D, solid solns. with aluminum arsenide
RL: USES (Uses)
(bipolar transistors contg. heterojunctions from gallium arsenide and, for temp. sensors)

IT 22831-42-1D, solid solns. with gallium arsenide
RL: DEV (Device component use); USES (Uses)
(bipolar transistors contg. heterojunctions from gallium arsenide and, for temp. sensors)

IT 1303-00-0, uses and miscellaneous
RL: DEV (Device component use); USES (Uses)
(bipolar transistors contg., for temp. sensors)

IT 101964-86-7
RL: TEM (Technical or engineered material use); USES (Uses)
(diffusion barriers from, in heterojunction bipolar transistors)

IT 7440-57-5, properties
RL: TEM (Technical or engineered material use); USES (Uses)
(diffusion barriers of iridium-tantalum alloy for, in heterojunction bipolar transistors)

L42 ANSWER 1 OF 1 HCAPLUS COPYRIGHT 2002 ACS
AN 1997:475876 HCAPLUS
DN 127:89684
TI Magnetic devices and magnetic sensors using thereof
IN Mizushima, Koichi; Konno, Teruyuki; Inomata, Koichiro; Yamauchi, Hisashi
PA Toshiba Corp., Japan
SO Jpn. Kokai Tokkyo Koho, 16 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 09128719	A2	19970516	JP 1996-189366	19960718
	JP 3217703	B2	20011015		

PRAI JP 1995-225625 A 19950901

AB The title magnetic sensors have a three terminal device comprising an **emitter**, a base, and a collector, wherein the semiconductor collector layer and a magnetic laminated base film make a Schottky junction. The magnetic laminated base film has an nonmagnetic film bound between opposing magnetic films. The metallic **emitter** film and base film are connected each other via a **tunneling** insulator **film**. The sensors provide variation of current across the magnetic device by magnetization orientation of the magnetic film changed by an external magnetic field direction. The devices gives high sensitivity by low c.d.

L45 ANSWER 1 OF 44 HCAPLUS COPYRIGHT 2002 ACS
 AN 2001:923312 HCAPLUS
 DN 136:46996
 TI Manufacture of cathodes for electron tubes
 IN Chiba, Toru
 PA Nec Corp., Japan
 SO Jpn. Kokai Tokkyo Koho, 5 pp.
 CODEN: JKXXAF
 DT Patent
 LA Japanese
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 2001351505	A2	20011221	JP 2000-169566	20000606
AB	Ni powders, emitter material powders and rare earth oxide powders are mixed, and hot pressed to form sintered bodies, the bodies are processed into cathode pellets, and Ni-Cr alloy films are formed on the other side of electron-emitting surface. Cr is fast and uniformly diffused in the pellets so that manufd. cathodes have stable electron emission characteristic.				

L45 ANSWER 2 OF 44 HCAPLUS COPYRIGHT 2002 ACS
 AN 2001:381858 HCAPLUS
 DN 135:124859
 TI Antireflection coated refractory metal matched **emitters** for use with GaSb thermophotovoltaic generators
 AU Fraas, Lewis; Samaras, John; Avery, James; Minkin, Leonid
 CS JX Crystals Inc, Issaquah, WA, 98027, USA
 SO Conference Record of the IEEE Photovoltaic Specialists Conference (2000), 28th, 1020-1023
 CODEN: CRCNDP; ISSN: 0160-8371

PB Institute of Electrical and Electronics Engineers
 DT Journal
 LA English
 AB GaSb thermophotovoltaic cells can be combined with IR **emitters** to produce elec. power. In this application, both power d. and efficiency are important. High power d. requires a practical target **emitter** temp. of 1600.degree. K. In order to reach this temp., spectral efficiency becomes extremely important. Radiation with wavelengths greater than 1.8 .mu.m cannot be converted by the GaSb cells; instead, this long wavelength radiation overheats the cells, limiting power d. and efficiency. A soln. is to use refractory-metal coated **emitters**, because metals have low emittance at long wavelengths. Further, an antireflection (AR) coating on the metal can enhance the emittance in the cell convertible band. A spectral efficiency of 75% has been demonstrated for an AR coated **emitter** and a GaSb cell power d. of 1.5 W/cm² has been measured with an AR coated tungsten **emitter** operating at 1555.degree. K.

L45 ANSWER 3 OF 44 HCAPLUS COPYRIGHT 2002 ACS
 AN 2001:323080 HCAPLUS
 DN 135:85335
 TI Development of thin-film **metal** hydrides for integration into field emission displays
 AU Chalamala, Babu R.; Reuss, Robert H.
 CS Motorola, Inc., Semiconductor Products Sector, Digital DNA Laboratory, Tempe, AZ, 85284, USA
 SO Applied Physics Letters (2001), 78(19), 2967-2969

CODEN: APPLAB; ISSN: 0003-6951
 PB American Institute of Physics
 DT Journal
 LA English
 AB We report on the development of solid-state hydrogen sources utilizing thin-film metal hydrides. We demonstrate that integration of these metal hydride thin films facilitate a practical method to introduce controlled amounts of hydrogen into sealed field emission display assemblies. To prove the concept, we operated Mo field emitter arrays without emission current loss for 400 h of continuous operation with titanium-hydride-coated stainless steel anode plates. Comparable arrays operated in the absence of hydride films, but in ultrahigh vacuum, had emission current degradation of over 50% in less than 100 h of operation.

RE.CNT 13 THERE ARE 13 CITED REFERENCES AVAILABLE FOR THIS RECORD
 ALL CITATIONS AVAILABLE IN THE RE FORMAT

L45 ANSWER 4 OF 44 HCAPLUS COPYRIGHT 2002 ACS
 AN 2000:574048 HCAPLUS
 DN 133:153231
 TI Antireflection coated refractory metal matched IR emitter for use in thermophotovoltaic generators
 IN Fraas, Lewis M.; Magendanz, Galen; Avery, James E.
 PA Jx Crystals Inc., USA
 SO PCT Int. Appl., 21 pp.
 CODEN: PIXXD2
 DT Patent
 LA English
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2000048231	A2	20000817	WO 1999-US24736	19991022
	W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
	RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
	AU 2000051216	A5	20000829	AU 2000-51216	19991022
PRAI	US 1998-113353P	P	19981221		
	US 1999-120817P	P	19990219		
	US 1999-406727	A	19990928		
	WO 1999-US24736	W	19991022		

AB Thermophotovoltaic (TPV) elec. power generators have emitters with IR outputs matched with usable wavelengths for converter cells. The emitters have durable substrates, optional refractory isolating layers, conductive refractory metal or inter-metallic emitter layers, and refractory metal oxide antireflection layers. SiC substrates have W or TaSi2 emitter layers and 0.14 .mu. ZrO2 or Al2O3 antireflection layers used as IR emitters for GaSb converter cells in TPV generators.

L45 ANSWER 5 OF 44 HCAPLUS COPYRIGHT 2002 ACS
 AN 2000:541583 HCAPLUS
 DN 133:289670
 TI Studies on the interaction between thin film materials and Mo field

emitter arrays
AU Chalamala, Babu R.; Reuss, Robert H.
CS Flat Panel Display Division, Motorola Incorporated, Tempe, AZ, 85284, USA
SO Journal of Vacuum Science & Technology, B: Microelectronics and Nanometer
Structures (2000), 18(4), 1825-1832
CODEN: JVTBD9; ISSN: 0734-211X
PB American Institute of Physics
DT Journal
LA English
AB A simple method for the evaluation of materials suitable for the
fabrication of field emission vacuum microelectronic devices is presented.
Since there can be a wide range of electron and ion interactions with the
device, it is important to be able to quickly assess if a material may
have a particular adverse effect on emission performance under operational
conditions. The technique is based on the sensitivity of a large field
emitter array to the outgassing or desorption of gas species from
thin films under electron beam excitation. Mo field **emitter**
arrays degraded rapidly with stainless steel anodes coated with various
oxide materials. The extent of degrdn. is the most rapid with SiO₂,
Si₃N₄, and MoO₃ thin films. Stainless steel anodes with Mo and Nb thin
films show a faster degrdn. rate than stainless steel anodes, most likely
because of native oxides grown during processing and handling. The
emission behavior in the presence of Ir, Pd, Al, Zn, and Ti **metal**
films and barrier materials like C and TaN is similar to stainless
steel ref. data. Once the oxide films are covered with barrier layers
like C and TaN, emission decay rates approach the values obtained with
stainless steel ref. anodes. The obsd. emission current degrdn. is
consistent with a model based on the liberation of oxygen from the surface
of electron beam bombarded materials. Using controlled oxygen exposure
expts., the authors detd. the equiv. local oxygen pressures in the
presence of various thin films. With thin films of Nb, ZrO₂, Ta₂O₅, MgO,
Nb₂O₅, and Al₂O₃, the emission degrdn. is akin to having a local O₂
partial pressure in the 1 .times. 10⁻⁷-1 .times. 10⁻⁶ Torr range and with
Mo, MoO₃, Si₃N₄, and SiO₂, this is equiv. to having local O₂ pressures of
1 .times. 10⁻⁵ Torr.

RE.CNT 20 THERE ARE 20 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L45 ANSWER 6 OF 44 HCAPLUS COPYRIGHT 2002 ACS
AN 2000:537534 HCAPLUS
DN 133:113541
TI P-type ohmic contact forming method
IN Park, Sung-ho; Park, Chol-soon; Park, Hyung-moo
PA Korea Electronics + Telecommunication Research Institute, S. Korea
SO Repub. Korea, No pp. given
CODEN: KRXXFC
DT Patent
LA Korean
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	KR 9706734	B1	19970429	KR 1993-27627	19931214
AB	A technique is described that provides a method for forming a p-type ohmic contact of a heterojunction bipolar transistor. The method includes the steps of (A) sequentially forming a buffer layer, subcollector layer, collector, base layer, spacer layer, emitter layer and capping layer on a semi-insulating AlGaAs substrate, (B) depositing an emitter metal on the capping layer and then mesa-etching the capping layer to the surface of the base layer using the emitter				

metal as a mask, (C) forming a p-type base metal by sequentially depositing a 1st Cr-metal layer, a AuZn alloy layer, 2nd Cr metal layer, Pd metal layer and Au metal layer on the etched base layer and (D) continuously covering the entire substrate with a Si oxide layer and Si nitride layer. Thereby, it is possible to improve the characteristic of the p-type ohmic contact.

L45 ANSWER 7 OF 44 HCAPLUS COPYRIGHT 2002 ACS

AN 2000:50029 HCAPLUS

DN 132:115264

TI Electron **emitter**, electron source, imaging device, and manufacture of the **emitter**, the source, and the device

IN Kobayashi, Tamaki

PA Canon K. K., Japan

SO Jpn. Kokai Tokkyo Koho, 19 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI	JP 2000021290	A2	20000121	JP 1998-187978 19980703
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AB The electron **emitter** consists of a pair of electrode and an elec. conductor film between the electrodes, which is prep'd. by (1) forming a film comprising 2 different metal layers (e.g., the top Pd layer and the bottom Ta or W layer), and (2) performing the so-called forming, i.e., supplying elec. current to the film and forming a narrow crack which emits electrons. The 2 layers are alloyed at the crack in the 2nd step and electrons are efficiently emitted from the crack which is prevented from excessive expansion. An electron source or an imaging device, e.g., electroluminescent display, etc., is manufd. by combining plurality of the electron **emitters**.

L45 ANSWER 8 OF 44 HCAPLUS COPYRIGHT 2002 ACS

AN 1999:690271 HCAPLUS

DN 131:304539

TI Field-emission cold cathode and manufacture of the cathode

IN Muroda, Masao

PA Nippon Denshi K. K., Japan

SO Jpn. Kokai Tokkyo Koho, 9 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI	JP 11297189	A2	19991029	JP 1998-99391 19980410
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AB The cathode has an **emitter** at the center of a polycryst. metal filament and the **emitter** comprises a body-centered cubic crystal and 2 face-centered cubic crystal layers which successively cover the bcc crystal. The cathode is manufd. by successively forming the 2 fcc layers on the bcc metal so that a nanoemitter with ternary alloy surface layer (comprising the 3 crystals) is formed. The **emitter** having the ternary alloy surface film has merits compared with conventional ones, e.g., self regeneration effect by heat diffusion on the surface, etc.

L45 ANSWER 9 OF 44 HCAPLUS COPYRIGHT 2002 ACS

AN 1999:639279 HCAPLUS

DN 131:304368
 TI The mechanism of localized in-pile corrosion of zirconium alloys. (II)
 AU Ozaki, Satoru
 CS Nuclear Engineering Laboratory, Toshiba Corporation, Shinsugita-cho,
 Isogo-ku, Yokohama, 235-8523, Japan
 SO J. Nucl. Sci. Technol. (1999), 36(7), 605-612
 CODEN: JNSTAX; ISSN: 0022-3131
 PB Atomic Energy Society of Japan
 DT Journal
 LA English
 AB The ".beta.-induced elec. fields" in a reactor, proposed in a previous
 paper, are estd. more quant. in two cases. One of the cases is that of
 the shadow formation on the surface of Zircaloy fuel channels near
 stainless steel control blades in BWRs by adapting a multi-element model
 analogous to an elec. circuit instead of the preliminary model in the
 previous paper. The other is that of the Halden reactor irradn. expts.,
 in which the localized in-pile corrosion of Zircaloy fuel claddings near
 platinum was reported by Y. Etoh et al. (1997), by adapting the detailed
 irradn. data in the report. As a result, both cases are explained by two
 parallel mechanisms affected by the ".beta.-induced elec. fields" on and
 in zirconium oxide and high-purity water. One of the mechanisms is the
 formation of localized electrochem. cells which occur between the surface
 of zirconium oxide adjacent to bare and grounded metals
 such as stainless steel or platinum and its neighboring parts. The other
 is the retardation of oxygen ions in the zirconium oxide by the space
 charge effect, proposed by S. Nanikawa et al. (1998), according to the
 level of the induced elec. potential.

RE.CNT 11 THERE ARE 11 CITED REFERENCES AVAILABLE FOR THIS RECORD
 ALL CITATIONS AVAILABLE IN THE RE FORMAT

L45 ANSWER 10 OF 44 HCAPLUS COPYRIGHT 2002 ACS
 AN 1999:307728 HCAPLUS
 DN 131:133621
 TI Mineralogy, environment, and nickel processing. The international Kola
 project
 AU Gregurek, Dean; Reimann, Clemens; Stumpf, Eugen F.
 CS Inst. Geowissenschaften, Abteilung Mineralogie Petrologie, Montanuniv.,
 Leoben, A-8700, Austria
 SO Berg- Huettenmaenn. Monatsh. (1999), 144(4), 146-151
 CODEN: BHMMAM; ISSN: 0005-8912
 PB Springer-Verlag Wien
 DT Journal
 LA German
 AB The mineralogical compn. of snow filtrates and soil samples taken in the
 vicinity of Ni-Cu mines and smelters in the Kola region (Russia) were
 investigated within the context of an international project to det. the
 extent of the environmental impact of the Ni processing industry. Annual
 emissions which include 296,000 t of SO₂ and significant amt. of metals
 have led to almost complete destruction of vegetation and severe damage to
 soils in the vicinity of individual emitters. Samples from 15
 selected sites nearby mines and smelters were prep'd. for microscopical and
 electron microprobe anal. Several minerals (sulfides, oxides, and
 silicates) and technogenic phases (Cu-Ni-Fe-Co sulfides globules and
 slags) were identified in the polished specimens. Based on their compn.,
 they can be attributed to specific emitters and to certain
 metallurgical processes. It is discussed that the lack of mineralogical
 data can limit the significance of chem. analyses of heavy metal
 processing polluted areas for the hazard assessment of heavy metal
 mobility in the soil.

RE.CNT 18 THERE ARE 18 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L45 ANSWER 11 OF 44 HCAPLUS COPYRIGHT 2002 ACS

AN 1999:175543 HCAPLUS

DN 130:198785

TI Rare earth **emitter**

IN Sarraf, David B.

PA Thermal Corp., USA

SO U.S., 4 pp.

CODEN: USXXAM

DT Patent

LA English

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI US 5879473	A	19990309	US 1997-791375	19970130
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AB The app. is a selective radiation **emitter** to be used in conjunction with a photocell for thermophotovoltaic generation of electricity from heat. One embodiment of the **emitter** is a layer of selectively radiating rare earth **oxide** bonded onto a heated base **metal** layer by the use of an intermediate thin layer of porous metal powder. Another embodiment is an **emitter** of a thick metal powder layer combined with a rare earth oxide which is formed into the voids in the metal powder structure and bonded to a metal substrate which is to be heated. In this embodiment the metal powder grains are gold plated to limit their emissivity, so that emission outside the desired band of the rare earth oxide is greatly reduced.

RE.CNT 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD

ALL CITATIONS AVAILABLE IN THE RE FORMAT

L45 ANSWER 12 OF 44 HCAPLUS COPYRIGHT 2002 ACS

AN 1998:779690 HCAPLUS

DN 130:46219

TI Fabrication of compound semiconductor device and compound semiconductor device itself

IN Asano, Tetsuo

PA Sanyo Electric Co., Ltd., Japan

SO Jpn. Kokai Tokkyo Koho, 15 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI JP 10321642	A2	19981204	JP 1997-128548	19970519
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AB The title method involves etching cap and **emitter** layers using a dummy **emitter** as a mask to remove the cap layer and a portion of the **emitter** layer, forming a side wall protective layer, etching the remaining **emitter** layer using the dummy **emitter** layer and side wall protective layer as a mask to form an **emitter** ledge, forming a resist film, etching back the resist film to expose the dummy **emitter** layer and side wall protective layer, wet etching the dummy **emitter** layer and side wall protective layer to expose the remaining cap layer, depositing a 1st **metal** film on the overall surfaces, and lifting off the resist film to selectively form an **emitter** electrode from the 1st **metal** film on the cap layer. Addnl., the method involves forming a collector electrode from a 2nd **metal** film.

Specifically, the 1st and 2nd metal films may comprise Pt/Ti/Pt/Au and AuGe/Ni, resp. By wet etching, damaging of other regions is prevented, and by lifting off, an isolated electrode is formed without ion milling. A compd. semiconductor device fabricated by the above method is also described.

L45 ANSWER 13 OF 44 HCPLUS COPYRIGHT 2002 ACS
 AN 1997:480696 HCPLUS
 DN 127:103113
 TI Electron-emitting element and manufacture thereof
 IN Gamo, Hidenorori; Kanamaru, Seigo; Ito, Junji
 PA Toppan Printing Co., Ltd., Japan; Agency of Industrial Sciences and Technology
 SO Jpn. Kokai Tokkyo Koho, 12 pp.
 CODEN: JKXXAF
 DT Patent
 LA Japanese
 FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI JP 09129123	A2	19970516	JP 1996-90337	19960319
JP 3012517	B2	20000221		

PRAI JP 1995-138750 A 19950511
 JP 1995-240999 A 19950825

AB In the electron-emitting element comprising a substrate, an emitter wiring layer, an insulating layer and a gate electrode laminated in order, a opening section arranged in the gate electrode and the insulating layer for reaching the emitter wiring layer, in the opening section, a cone-shaped emitter formed on the emitter wiring layer and without contact with the gate emitter, the emitter wiring layer is formed from a metal (e.g., Cr, Cr-Al alloy) thin film, and the emitter is formed from non-single cryst. Si. The element is esp. useful for field emitter array (FEA) in flat display device.

L45 ANSWER 14 OF 44 HCPLUS COPYRIGHT 2002 ACS
 AN 1997:475876 HCPLUS
 DN 127:89684
 TI Magnetic devices and magnetic sensors using thereof
 IN Mizushima, Koichi; Konno, Teruyuki; Inomata, Koichiro; Yamauchi, Hisashi
 PA Toshiba Corp., Japan
 SO Jpn. Kokai Tokkyo Koho, 16 pp.
 CODEN: JKXXAF
 DT Patent
 LA Japanese
 FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI JP 09128719	A2	19970516	JP 1996-189366	19960718
JP 3217703	B2	20011015		

PRAI JP 1995-225625 A 19950901

AB The title magnetic sensors have a three terminal device comprising an emitter, a base, and a collector, wherein the semiconductor collector layer and a magnetic laminated base film make a Schottky junction. The magnetic laminated base film has an nonmagnetic film bound between opposing magnetic films. The metallic emitter film and base film are connected each other via a tunneling insulator film. The sensors provide variation of current across the magnetic device by magnetization orientation of the magnetic film

changed by an external magnetic field direction. The devices gives high sensitivity by low c.d.

L45 ANSWER 15 OF 44 HCPLUS COPYRIGHT 2002 ACS
 AN 1997:134688 HCPLUS
 DN 126:151612
 TI Integrated circuits and their manufacture
 IN Nishihara, Shinji; Ikeda, Shuji; Ishida, Shinichi; Suzuki, Masayasu; Asayama, Masaichiro
 PA Hitachi Ltd, Japan
 SO Jpn. Kokai Tokkyo Koho, 10 pp.
 CODEN: JKXXAF
 DT Patent
 LA Japanese
 FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 08330458	A2	19961213	JP 1995-136254	19950602

PI AB The circuit is composed of MISFETs and bipolar transistors on the main surface of a semiconductor substrate, and has an **emitter** electrode from a doped polycryst. Si-refractory metal silicide laminate on the **emitter** region, a 1st interlayer insulating film contg. a borophosphosilicate or phosphosilicate glass film, and wiring consisting of refractory metal and conductor films, connected to the **emitter** electrode and the source or the drain electrode of the MISFET through contact holes. A low-resistance refractory metal silicide is formed at the interface between the refractory metal silicide of the **emitter** electrode and the refractory metal film in annealing, hence consumption of Si produces more holes and activates the impurity in the polycryst. Si film; thus the impurity concn. in the **emitter** region is raised and current amplification of **emitter** grounding is improved without deep diffusion of the impurity in the source-drain region; the contact resistance of the source or the drain is lowered by formation of low-resistance refractory metal silicide, and characteristics of the **emitter** electrode are improved as recombination centers of holes in the polycryst. Si film disappear.

L45 ANSWER 16 OF 44 HCPLUS COPYRIGHT 2002 ACS
 AN 1996:237597 HCPLUS
 DN 124:275978
 TI Memory and amplifier devices
 IN Sakakima, Hiroshi; Irie, Yasusuke; Kawawake, Yasuhiro; Satomi, Mitsuo
 PA Matsushita Electric Ind Co Ltd, Japan
 SO Jpn. Kokai Tokkyo Koho, 5 pp.
 CODEN: JKXXAF
 DT Patent
 LA Japanese
 FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 08018118	A2	19960119	JP 1994-149230	19940630

PI AB The memory device is constructed from (1) a laminate conductor which consists of a hard magnetic film (e.g., from Co-Pt) having a good squareness in hysteresis and large variation of magnetoresistance with minute change of magnetic field, a soft magnetic film (e.g., from NixCoyFez with x = 0.6-0.9, yr = 0.05-0.4, and z = 0-0.3 or x = 0-0.4, yr = 0.2-0.5, and z = 0-0.5) easy in magnetic inversion by a change in a weak magnetic field, and a nonmagnetic metal film (e.g.,

Cu, Ag, or Au) therebetween for lowering of magnetic bonding between the magnetic films and (2) a conductive lead located in the vicinity of the laminate conductor as sepd. from the laminate conductor by an insulating film, forming the magnetic films to an **emitter** and a collector, or vice versa, and the nonmagnetic film to the base, with placement of a voltage detector or a load on the side of the collector. The amplifier device has the same construction.

L45 ANSWER 17 OF 44 HCPLUS COPYRIGHT 2002 ACS

AN 1996:231786 HCPLUS

DN 124:304832

TI Electron-emitting device

IN Sano, Kenya; Kawakubo, Takashi

PA Tokyo Shibaura Electric Co, Japan

SO Jpn. Kokai Tokkyo Koho, 5 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI JP 08031302 A2 19960202 JP 1994-165164 19940718

AB The device comprises an insulating film of an **oxide** of Ta-contg. amorphous **alloy** sandwiched between a pair of electrodes.

Preferably, the amorphous alloy contains Ta and Al as main components.

The device generates low diode current, and effectively emits electrons.

L45 ANSWER 18 OF 44 HCPLUS COPYRIGHT 2002 ACS

AN 1995:532014 HCPLUS

DN 122:280100

TI Inorganic **emitter** materials and discharge electrodes for lamps

IN Takegawa, Yoshinobu; Sakon, Shigetoshi; Imamura, Hiroshi

PA Matsushita Electric Works Ltd, Japan

SO Jpn. Kokai Tokkyo Koho, 5 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI JP 06223774 A2 19940812 JP 1993-9544 19930122

AB The electrodes are prep'd. by depositing on **metal** wires successively with **metal oxides** and inorg. materials contg. **emitter** and reducing materials. The **emitter** materials may be alk. earth carbonates. The reducing material may be a Group IIIA, IVA, or VA metal. The **metal oxide** may be alumina.

L45 ANSWER 19 OF 44 HCPLUS COPYRIGHT 2002 ACS

AN 1995:394725 HCPLUS

DN 122:149563

TI Electrode in fluorescent lamps

IN Takegawa, Yoshinobu; Sakon, Shigetoshi; Yamada, Shuji

PA Matsushita Electric Works Ltd, Japan

SO Jpn. Kokai Tokkyo Koho, 5 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 06150881	A2	19940531	JP 1992-300164	19921110
AB	In manuf. of fluorescent lamp electrode having aluminum oxide covered metal wire and emitter material coated on the aluminum oxide layer, the surface of the aluminum oxide layer is prep'd. in whisker-like shape to ensure high adhesive intensity of the emitter material to the ion bombardment. The metal wire is made of Fe-Cr-Al alloy of compn. ranges: Cr 10 - 30 wt%, Al 1 - 10 wt% and Fe for the rest.				

L45 ANSWER 20 OF 44 HCPLUS COPYRIGHT 2002 ACS
 AN 1994:591522 HCPLUS
 DN 121:191522
 TI Electron emitting elements, light emitting elements, and image display devices
 IN Kaneko, Tetsuya; Nomura, Ichiro; Suzuki, Hidetoshi; Nakamura, Naohito; Iwai, Hisami; Takeda, Toshihiko
 PA Canon Kk, Japan
 SO Jpn. Kokai Tokkyo Koho, 20 pp.
 CODEN: JKXXAF
 DT Patent
 LA Japanese
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 06020590	A2	19940128	JP 1992-194564	19920630
AB	JP 3023734 B2 20000321 An electron emitting-element consists of a thermal fusion bonded vacuum container, a metal electrode, and an electron emitting base; the surface of the electrode consists of an alloy of the electrode metal and the elec. cond. of the alloy oxide is greater than that of the oxide of the electrode metal. An image display device is manufd. from the electron emitting elements, which are arranged on an electron source, and a face plate, and a light emitting device is manufd. by placing a fluorescing element and a face plate on an array of electron emitting devices. The electrode metal is Ni and the surface is a Ni alloy contg. 5-20 mol.% Ag or Li, or Cr and the surface is a Cr alloy contg. 0.35-1.4 mol.% Ti or 1.25-5 mol.% Ni.				

L45 ANSWER 21 OF 44 HCPLUS COPYRIGHT 2002 ACS
 AN 1994:259246 HCPLUS
 DN 120:259246
 TI Impregnated cathodes for cathode-ray tubes and manufacture thereof
 IN Taguchi, Tadanori; Suzuki, Yukio; Saito, Shunji
 PA Hitachi Ltd, Japan
 SO Jpn. Kokai Tokkyo Koho, 7 pp.
 CODEN: JKXXAF
 DT Patent
 LA Japanese
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 05250981	A2	19930928	JP 1992-48709	19920305
AB	The cathode consists of refractory metal sections from Mo and/or W, and electron emitter sections from compds. contg. at least Ba and O, and W, Sc, and O, and is prep'd. by mixing of pellets from a powder of W and/or Mo, the compd. contg. Ba and O, and that contg. W, Sc, and O, pelletization of the mixt., debinderization of the pellets in a H2 atm.,				

sintering of W and/or Mo and fixation of **emitters** thereon, and assembly with a barrier layer into a sleeve.

L45 ANSWER 22 OF 44 HCAPLUS COPYRIGHT 2002 ACS
 AN 1994:122837 HCAPLUS
 DN 120:122837
 TI Electron **emitters** and manufacturing thereof
 IN Yoshida, Yoshihiro; Kozuka, Takeshi; Kobayashi, Hiroshi
 PA Ricoh Kk, Japan
 SO Jpn. Kokai Tokkyo Koho, 7 pp.
 CODEN: JKXXAF
 DT Patent
 LA Japanese
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI	JP 05198252	A2	19930806	JP 1992-7997	19920121
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AB The title manufg. involves depositing a **metal** film on an elec. conductive layer, anodizing the **metal** film to form an insulator **film** having micropore through-holes, selectively etching the insulator film to give contact holes, and depositing electron emitting material in the contact holes. The arrangement gives easy application for large scale **emitters** in economical manufg.

L45 ANSWER 23 OF 44 HCAPLUS COPYRIGHT 2002 ACS
 AN 1994:93158 HCAPLUS
 DN 120:93158
 TI Implant-free heterojunction bipolar transistor integrated circuit fabrication process
 IN Prasad, Jayasimha S.; Park, Song W.; Vetanen, William A.; Beers, Irene G.; Haynes, Curtis M.
 PA Tektronix, Inc., USA
 SO U.S., 12 pp.
 CODEN: USXXAM
 DT Patent
 LA English
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI	US 5268315	A	19931207	US 1992-940588	19920904
	GB 2270418	A1	19940309	GB 1993-17381	19930820
	DE 4329189	A1	19940310	DE 1993-4329189	19930830
	JP 06232150	A2	19940819	JP 1993-244000	19930903

PRAI US 1992-940588 19920904
 AB The process can fabricate n-p-n heterojunction bipolar transistors, Schottky diodes, MIM capacitors, spiral inductors, and NiCr resistors. Two levels of interconnect metal are available. The first level **metal** is a conventional **dielec.-insulated metal** conductor. The second level metal includes an air-bridge for contacting the HBT **emitters**, which are on top of three level mesa structures. It is also an advanced low loss, low capacitance, air dielec. conductor useful for long interconnects and inductors. MIM capacitors are formed by sandwiching silicon nitride between the first layer metal and a capacitor top plate made with landed air-bridge metal. Precision thin film resistors are fabricated by depositing NiCr on silicon nitride. The three-level active mesa structure is etched down to the GaAs substrate, for lateral device isolation, with a truncated pyramidal shape which permits good step coverage of dielec. and metalization layers. The wet

etching process uses a compn. of H₃PO₄:H₂O₂:H₂O in a preferred ratio of about 3:1:25 for the AlGaAs/GaAs system.

L45 ANSWER 24 OF 44 HCAPLUS COPYRIGHT 2002 ACS
 AN 1993:683880 HCAPLUS
 DN 119:283880
 TI Noble-metal alloy composites for durable secondary electron emitters
 AU Aitov, R. D.; Bondarenko, G. G.; Korzhavyi, A. P.
 CS Moscow, Russia
 SO Metally (1993), (4), 223-5
 CODEN: MEALET
 DT Journal
 LA Russian
 AB The use was studied of Pt, Pd, and Ag composites with oxides for secondary electron emitters. The effect was studied of the surface charge on the electron emission. The best sources are obtained using Li₂O.

L45 ANSWER 25 OF 44 HCAPLUS COPYRIGHT 2002 ACS
 AN 1993:639441 HCAPLUS
 DN 119:239441
 TI Manufacture of heterojunction bipolar transistors
 IN Mitani, Katsuhiko; Masuda, Hiroshi; Mochizuki, Kazuhiro; Kusano, Chushiro
 PA Hitachi Ltd, Japan
 SO Jpn. Kokai Tokkyo Koho, 6 pp.
 CODEN: JKXXAF
 DT Patent
 LA Japanese
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 05067628	A2	19930319	JP 1991-230042	19910910
AB	Manuf. of a heterojunction bipolar transistor includes: (a) successively forming a 1st-cond. high-concn. subcollector layer, a 1st-cond. low-concn. collector layer, a 2nd-cond. high-concn. base layer, a 1st-cond. emitter layer with a greater bandgap than the base layer, and a 1st-cond. high-concn. subemitter layer; (b) deposition of a 1st metal film (e.g., WSi) and a 1st insulator film on the high-concn. subemitter layer; (c) removing the metal film, the 1st insulator film, the high-concn. subemitter-forming region to create an emitter mesa; (d) surrounding the emitter mesa with a 2nd insulator film formed up to the height of the 1st insulator film; (e) removing the 1st insulator film exposed in the 2nd insulator film to create an opening; (f) forming a 2nd metal film (e.g., W) with a wider top surface than the bottom surface on the 1st metal film exposed in the opening; (g) removing the 2nd insulator film; and (h) forming a 3rd metal film (e.g., AuZn base electrode) on the 2nd metal film and on the base layer around the emitter mesa. The heterojunction bipolar transistor has decreased external base resistance, and is capable of fast operation.				

L45 ANSWER 26 OF 44 HCAPLUS COPYRIGHT 2002 ACS
 AN 1993:615278 HCAPLUS
 DN 119:215278
 TI Contamination of liquid-metal indium ion emitters
 AU Sieber, Anton; Fehringer, Michael; Ruedenauer, Friedrich G.; Steiger, Wolfgang
 CS Hauptabte. Angew. Phys., Forschungszent. Seibersdorf, Switz.

SO Oesterr. Forschungszent. Seibersdorf, [Ber.] OEFZS (1993), OEFZS-4674, 23 pp.
 CODEN: OFSODK; ISSN: 0253-5270
 DT Report
 LA German
 AB Electron microprobe anal. of liq.-metal In ion **emitter** needles burning against a stainless steel collector showed that they are contaminated with high concns. of Fe and Cr with exception of the apex area, where the indium is clean. The contamination freeze in platelike structures under which In is flowing towards the apex. Due to breakage of the plates and reflow of In, the **metal film** on the needle is a multilayer film, in which In-rich layeres alternate with contamination-rich layers.

L45 ANSWER 27 OF 44 HCAPLUS COPYRIGHT 2002 ACS

AN 1993:263595 HCAPLUS

DN 118:263595

TI Alkali metal source for photomultiplier

IN Koike, Takashi; Inazuru, Tsutomu

PA Hamamatsu Photonics K. K., Japan

SO Jpn. Kokai Tokkyo Koho, 5 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI JP 04242040	A2	19920828	JP 1991-16067	19910114
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AB The source, which is mounted on a lead placed along the inner wall of a photomultiplier, comprises a pair of metal hats, each of which encases a pellet contg. reactants which produce an alkali metal upon heating and a heater plate, wherein the pellets are placed on both sides of the heater, the hats are placed thereon and sealed off, the heating generates the metal vapor which leaks through a heating-created gap in the seal, and the vapor converts the precoat on the dynodes to the photoelectron **emitter**.

L45 ANSWER 28 OF 44 HCAPLUS COPYRIGHT 2002 ACS

AN 1993:244726 HCAPLUS

DN 118:244726

TI Electron **emitter** and image-display and imaging devices using the **emitter**

IN Takimoto, Kiyoshi; Matsuda, Hiroshi; Kawagishi, Hideyuki; Morikawa, Yuko

PA Canon K. K., Japan

SO Jpn. Kokai Tokkyo Koho, 11 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI JP 04286828	A2	19921012	JP 1991-74209	19910315
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JP 3185060	B2	20010709		
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AB In an electron **emitter** having a structure of an insulator layer between a pair of electrodes, the insulator layer is formed on a smooth electrode having a surface roughness .1 toreq.1 nm , and the electrons are emitted from the surface opposite to the smooth electrode by applying an elec. potential between the electrodes. Specifically, the smooth electrode may comprise a noble metal on its **alloy**, and the

insulator film may comprise a Langmuir-Blodgett film. The elec.-field concn. is prevented and a uniform emission is obtained.

L45 ANSWER 29 OF 44 HCAPLUS COPYRIGHT 2002 ACS
AN 1993:85213 HCAPLUS

DN 118:85213

TI Corrosion-resistant iron alloy for **emitter** of far infrared

IN Ishii, Kazuhide; Kawasaki, Tatsuo

PA Kawasaki Steel Corp., Japan

SO Jpn. Kokai Tokkyo Koho, 3 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI JP 04173999	A2	19920622	JP 1990-295420	19901102
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AB The **emitters** for far-IR radiation are manufd. from Fe alloy strip contg. Cr 6-30, Al 2-8, and Zr 0.2-1.5%, and are pretreated for oxide coating with the thickness ≥ 0.2 mg/cm². The Fe-alloy strip is pretreated for >3 min at 900-1200.degree. in oxidizing atm. The **emitters** are useful for IR-heating app., esp. in heating of foods or drying of coatings.

L45 ANSWER 30 OF 44 HCAPLUS COPYRIGHT 2002 ACS

AN 1992:560481 HCAPLUS

DN 117:160481

TI Small-size fluorescent lamp

IN Tagawa, Koji; Uenari, Seiichi; Ikeuchi, Mitsuru; Onishi, Yasuo

PA Ushio Denki K. K., Japan

SO Jpn. Kokai Tokkyo Koho, 4 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI JP 04073853	A2	19920309	JP 1990-184167	19900713
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AB The lamp has 2 electrode substrates, each of which are coated with a getter and a Hg compd. on 1 of the surfaces and the other, resp., joined at $\angle 30$.degree. in opening angle, and an **emitter**-loaded metal mesh between the electrode substrates for a pair of opposing electrodes sealed with an inert gas at 80-130 torr and Hg in a glass tube ≤ 10 mm in inner diam., and is operated at ≤ 15 W in power consumption. The lamp has durability and suppression of blackening of the bulb due to scattering of the **emitter** material.

L45 ANSWER 31 OF 44 HCAPLUS COPYRIGHT 2002 ACS

AN 1992:265230 HCAPLUS

DN 116:265230

TI IR **emitters** with protected reflection layer and their preparation

IN Goebel, Wolfgang; Schmitz, Klaus; Wild, Wolfgang

PA Heraeus Quarzglas G.m.b.H., Germany

SO Ger. Offen., 3 pp.

CODEN: GWXXBX

DT Patent

LA German

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	DE 4021798	A1	19920206	DE 1990-4021798	19900709
	US 5276763	A	19940104	US 1992-949334	19920922
PRAI	DE 1990-4021798		19900709		
	US 1991-660489		19910225		
AB	IR sources comprising a quartz glass or quartz-based material housing contg. a heating element and provided with a metallic reflection film on the backside of the housing are described in which a protective layer of ZrO ₂ , SiO ₂ , SnO ₂ , or a mixt. of .gtrless.2 of these oxides is provided on the reflective layer. Prepn. of the sources includes applying a layer of .gtrless.1 thermally decomposable org. compd. of Zr, Sr, and/or Sn to the reflective layer and calcining at 600-950.degree..				

L45 ANSWER 32 OF 44 HCPLUS COPYRIGHT 2002 ACS
 AN 1988:612889 HCPLUS
 DN 109:212889
 TI An efficient and durable aluminum or aluminum alloy far-IR radiator
 IN Ishida, Shinichi; Yamada, Kikuo
 PA Nippon Aluminium Mfg. Co., Ltd., Japan
 SO Jpn. Kokai Tokkyo Koho, 6 pp.
 CODEN: JKXXAF
 DT Patent
 LA Japanese
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 63145797	A2	19880617	JP 1986-292121	19861208
AB	The radiator comprises an anodic oxide film including far-IR emitting materials. Optionally, the far-IR emitting material may comprise C, or an oxide of Fe, Cr, Ni, Co, Ti, Sn, Ag, Pb, Au, Mg, Mn, Zn, etc. The radiator is useful as a heating component.				

L45 ANSWER 33 OF 44 HCPLUS COPYRIGHT 2002 ACS
 AN 1983:604346 HCPLUS
 DN 99:204346
 TI Corrosion phenomena in metal-encapsulated tin-plated transistors
 AU Muecke, K. H.
 CS Fachber. Elektrotech. Maschinenbau, Fachhochsch. Landshut, Landshut, D-8300, Fed. Rep. Ger.
 SO Metalloberflaeche (1983), 37(10), 426-30
 CODEN: MOFEAV; ISSN: 0026-0797
 DT Journal
 LA German
 AB The corrosion in Ni encapsulated Sn-plated Si transistors was studied after > 10000 h operation at 40.degree.. Corrosion affects current-voltage properties. Corrosion occurs at the Au-plated base plate near the semiconductor, on the glass coating of the base plate, on the Al wires, and on the base and emitter region Al contacts. This corrosion is essentially due to electrodiffusion of ions on the glass and semiconductor and partly due to HCl or KCl etching, and anodization of the metal particles of the base plate. The corrosion is caused by H₂O produced by reaction of H from the Sn plate and O₂ trapped in the casing. It is prevented by using N₂ instead of air during the processing.

L45 ANSWER 34 OF 44 HCPLUS COPYRIGHT 2002 ACS
 AN 1980:536556 HCPLUS
 DN 93:136556

TI Cermet
 IN Clark, Grady Wayne; Holder, John Davis; Pasto, Arvid Eric
 PA United States Dept. of Energy, USA
 SO Ger. Offen., 20 pp.
 CODEN: GWXXBX

DT Patent
 LA German

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	DE 2947917	A1	19800604	DE 1979-2947917	19791128
	GB 2040908	A	19800903	GB 1979-32372	19790918
	GB 2040908	B2	19821027		
	JP 55076034	A2	19800607	JP 1979-154153	19791128

PRAI US 1978-964406 19781128

AB Directionally solidified cermets are obtained from a eutectic Mo-metal oxide by internal zone melting in a high-frequency elec. field. The induction heating parameters, elec. cond. of the cermet, and particle size are chosen to prevent melting of the external layers. The Mo content is 8-15 vo.%, and the balance is Cr2O3, LaCrO3, ZrO2-Al2O3, or ZrO2-CeO2. The cermets are useful for MHD electrodes, gas turbines, electron emitters, high-temp. valve seats, and cutting tools. The cermets are heated in the high-frequency field from room temp. to the eutectic melting temp. without radiative preheating. Thus, a mixt. of 150 g Cr2O3 -50/+80 mesh and 40 g Mo -100 mesh was hot pressed to 5 cm thick pellets in a Mo-lined graphite mold at 1550.degree. and 200 psig, and the pellets were heated in an induction field at 2.2 MHz from a 10 kW generator. The protective atm. was a CO-CO2 mixt. in a 10:1 vol. ratio. Output power of the generator was gradually increased to give a surface temp. of 1710.degree. (240.degree. below the eutectic temp.), and the pellets were moved through the inductor coil at 1 cm/h to obtain a directionally solidified Mo-Cr2O3 cermet.

L45 ANSWER 35 OF 44 HCPLUS COPYRIGHT 2002 ACS

AN 1980:190171 HCPLUS

DN 92:190171

TI Metal base for oxidic cathode

IN Ezawa, Masayoshi; Misumi, Akira; Shibata, Norio

PA Hitachi, Ltd., Japan

SO Jpn. Kokai Tokkyo Koho, 2 pp.

CODEN: JKXXAF

DT Patent

LA Japanese ,

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 54148138	A2	19791120	JP 1978-55585	19780512
	JP 61002748	B4	19860127		

AB The metal base used in the prepn. of oxide cathodes for electron tubes is obtained by heating a Ni base alloy contg. W and(or) Mo together with a min. amt. of a reducing metal (e.g. Zr) for 5-75 min at 80-150.degree. in a low-temp. O plasma, dissolving the W and(or) Mo oxide from the surface with an alkali and(or) H2O2 soln., and heating in vacuo (10-5-10-6 torr) for 5-60 min at 950-1100.degree. to form a dense surface structure. When an alkali-metal oxide which serves as an electron emitter is subsequently deposited, good adhesion takes place and spalling of the deposited electron-emitter layer does not occur readily.

L45 ANSWER 36 OF 44 HCPLUS COPYRIGHT 2002 ACS
 AN 1978:572615 HCPLUS
 DN 89:172615
 TI Thermal radiation **emitter** material
 IN Kuze, Takashi; Matsuki, Toshiharu; Nagaoka, Koji; Iwai, Naoji
 PA Tokyo Shibaura Electric Co., Ltd., Japan
 SO Japan. Kokai, 5 pp.
 CODEN: JKXXAF
 DT Patent
 LA Japanese
 FAN.CNT 3

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 53066836	A2	19780614	JP 1976-141832	19761126
	JP 58017265	B4	19830406		
	DE 2656167	A1	19770623	DE 1976-2656167	19761210

PRAI JP 1975-148912 19751212
 JP 1975-155379 19751224
 JP 1976-141832 19761126

AB A thermal radiation **emitter** useful in transmitter and receiver tubes is obtained by oxidizing a Cr-contg. alloy layer deposited on a substrate by heating in an oxidizing atm. Thus, a 5-.mu. film of an Fe **alloy** cong. V 0.2 and Cr 18% was sputtered on a 0.5-mm Fe plate, and the alloy coating was oxidized in moist H₂ (dew point 30.degree.) at 1200: for 1 h. The oxidized coating showed a thermal emissivity of 0.91%.

L45 ANSWER 37 OF 44 HCPLUS COPYRIGHT 2002 ACS
 AN 1977:494412 HCPLUS
 DN 87:94412
 TI Material and method of making secondary-electron **emitters**
 IN Henrich, Victor E.; Fan, John C. C.
 PA Massachusetts Institute of Technology, USA
 SO U.S., 6 pp.
 CODEN: USXXAM

DT Patent
 LA English
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 4038216	A	19770726	US 1975-604390	19750813
	US 4115228	A	19780919	US 1977-802203	19770531

PRAI US 1974-482109 19740624
 US 1975-604390 19750813

AB Efficient secondary-electron **emitters** can be obtained by using radio-frequency sputtering to codeposit a high-yield insulator (e.g. MgO) and a metal (e.g. Au, Ag, Pt) in the form of thick, finely grained cermet films and then differentially sputtering the film surface. The metal is sputtered away faster than the ceramic to leave a ceramic-rich surface layer having excellent secondary-emission properties for low-energy incident electrons. The presence of metallic particles in the bulk of the films and the small size of the ceramic particles greatly reduce surface charging while allowing the **emitter** film to be thick enough to have a long operating life under adverse device conditions. The cermet film consists of 50-90 vol.% MgO. The av. sizes of the MgO and the metal particles are chosen to allow charge tunneling from the surface of the MgO grains to the surrounding metal grains to eliminate surface charging. The surface of the cermet film has a compn. of 70-100 vol.% MgO.

L45 ANSWER 38 OF 44 HCPLUS COPYRIGHT 2002 ACS
 AN 1975:446688 HCPLUS
 DN 83:46688
 TI Rare earth **oxide-metal** eutectic composites
 AU Stendera, J. W.; Benzel, J. F.
 CS Sch. Ceram. Eng., Georgia Inst. Technol., Atlanta, Ga., USA
 SO J. Am. Ceram. Soc. (1975), 58(3-4), 116-19
 CODEN: JACTAW
 DT Journal
 LA English
 AB Unidirectionally solidified composites of a rare earth sesquioxide + CeO₂ + (Mo or W) and of CeO₂ + (Mo or W) were grown using the direct-radio-frequency-coupling internal-molten-zone technique. The dramatic fiber improvement on addn. of CeO₂ to the sesquioxide systems provides extra loosely bound O to increase the solv. of the **metal** in the molten **oxide**. The composites have extensive areas of ordered eutectic growth contg. 8-50 times 10⁶ metal fibers/cm²; the fibers are 0.1-0.6. μ diam. Fiber d. increased almost linearly as the growth rate was increased from 0.5 to .apprx.5.0 cm/hr. Slow growth favors the platelet metal morphology over the fiber morphology. Platelets are typically obsd. in areas where high impurity concns. are expected, e.g. at the top of the solidified zone and at grain boundaries. Selective etching of the metal fiber leaves a ceramic filter, and etching of the oxide exposes Mo pin tips for high-current d. field-effect electron **emitters**.

L45 ANSWER 39 OF 44 HCPLUS COPYRIGHT 2002 ACS
 AN 1974:542899 HCPLUS
 DN 81:142899
 TI Complex electron-optical study of molybdenum-platinum alloy thermionic **emitters**
 AU Shishkin, B. B.; Bakhtiyarov, R. S.
 CS Mosk. Gos. Univ. im. Lomonosova, Moscow, USSR
 SO Zh. Tekh. Fiz. (1974), 44(2), 387-99
 CODEN: ZTEFA3
 DT Journal
 LA Russian
 AB Studies of Mo-Pt thermionic **emitters** in emission and scanning electron microscopes and in an x-ray microanalyzer showed that the Pt forms a nonuniform **film** on the surface of the **alloy** which is not monat. and which leads to a nonuniform distribution of the work function over the surface. Equations were derived for the thermionic currents from **emitters** with nonuniform work functions and from those with complex conduction bands. Alloys of Mo with 1, 5, 10, and 15 wt. % Pt were studied.

L45 ANSWER 40 OF 44 HCPLUS COPYRIGHT 2002 ACS
 AN 1974:542897 HCPLUS
 DN 81:142897
 TI Preparation and properties of sputtered magnesia-gold, magnesia-silver, and magnesia-nickel cermet films
 AU Fan, John C. C.; Henrich, Victor E.
 CS Lincoln Lab., Massachusetts Inst. Technol., Lexington, Mass., USA
 SO J. Appl. Phys. (1974), 45(9), 3742-8
 CODEN: JAPIAU
 DT Journal
 LA English
 AB Finely grained films of 3 cermets-MgO/Au, MgO/Ag, and MgO/Ni-were grown by radio-frequency sputtering from composite targets. Electron-microscopic

studies show that MgO/Au and MgO/Ag films consist of small crystallites (usually <150 .ANG.) of both MgO and Au or Ag. In MgO/Ni films, it appears that amorphous Ni particles are embedded in a polycryst. MgO matrix. These cermet films are good secondary-electron emitters, especially in cases where differential-sputtering effects are large. In MgO/Au films, the Au particles sputter much faster than MgO resulting in a MgO-rich surface layer. In MgO/Ag films, the differential-sputtering effect is smaller, probably due to stronger interaction between MgO and Ag crystallites. In MgO/Ni, the Ni particles sputter slower than MgO resulting in a slightly Ni-rich surface (and hence in poorer electron-emission properties). The small particle sizes and the presence of metallic particles in the bulk of the films greatly reduce surface charging.

L45 ANSWER 41 OF 44 HCPLUS COPYRIGHT 2002 ACS
 AN 1974:430996 HCPLUS
 DN 81:30996
 TI Optimized photocathode for mobility measurements in liquids
 AU Martin, K.; Secker, P. E.
 CS Sch. Electron. Eng. Sci., Univ. Coll. North Wales, Bangor/Caerns, Wales
 SO J. Phys. E (1974), 7(6), 432-3
 CODEN: JPSIAE
 DT Journal
 LA English
 AB A description is given of the characteristics of a thin-film Au-Pd photosource arranged for back-illumination by uv photons. The emitter has a useful service life when operated in air or vapors of org. liqs. and exhibits a high photoelec. yield.

L45 ANSWER 42 OF 44 HCPLUS COPYRIGHT 2002 ACS
 AN 1974:407604 HCPLUS
 DN 81:7604
 TI High performance emitter for thermoelectronic diode
 IN Durand, Jean P.; Gillardeau, Jacques; Faron, Robert
 PA Commissariat a l'Energie Atomique
 SO Fr. Demande, 9 pp.
 CODEN: FRXXBL

DT Patent
 LA French
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	FR 2186706	A1	19740111	FR 1972-19985	19720602
	FR 2186706	B1	19741227		
	US 3883765	A	19750513	US 1973-363867	19730525
	DE 2328066	A1	19731213	DE 1973-2328066	19730601

PRAI FR 1972-19985 19720602
 AB A fissionable fuel, such as enriched U oxide, is encased in an emitter layer of at least 1 refractory metal. A preferred case, epitaxial and free of pores, obtained at .gtoreq.1200.degree. by the interaction of H at <1 mm pressure and the metal halogenide, preferably the hexafluoride, comprises a layer of Mo, followed by Mo-W, W, and a thin W surface layer oriented in the (110) plane. Deposition of the latter, the densest, most stable, and most suitable material for thermoelectronic emission, is enhanced by a temp. above 1500.degree. and the addn. of Cl or O to the gas mixt. Design details for improved mech. strength are described.

L45 ANSWER 43 OF 44 HCPLUS COPYRIGHT 2002 ACS

AN 1973:459191 HCAPLUS
 DN 79:59191
 TI Monolithic bipolar semiconductor device employing a cermet for both Schottky barrier and ohmic contact
 IN Breuer, David R.; Buie, James L.
 PA TRW Inc.
 SO U.S., 5 pp.
 CODEN: USXXAM
 DT Patent
 LA English
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 3737742	A	19730605	US 1971-185137	19710930

AB A cermet material is used in an integrated bipolar-transistor circuit to serve the following 4 functions simultaneously: (1) to contact n+ **emitter** and p+ base regions, (2) to form a Schottky barrier with a lightly doped n collector region, (3) to provide a barrier against interaction of Si with contact **metal**, and (4) to make thin-film resistors. The cermet film is deposited in a single flash-evapn. step to provide all 4 functions. It consists, e.g., of Cr 58 and SiO 42 wt. %, is 300 .ANG. thick, and has a sp. resistivity of 3 .times. 10-3 .OMEGA.-cm. The layer may be composed of Cr 50-65 with SiO 35-50 wt. % or Cr 40-60 with Si 40-60 wt. %. The Schottky barrier diode is in shunt with the base-collector junction, the base serving as the diode anode and the collector as the cathode, improving the high-speed performance of the transistor. A layer of contact metal over the cermet coating forms a common contact to the collector and base regions and sep. contacts to the **emitter** region and collector contact area. The circuit element is a digital correlator.

L45 ANSWER 44 OF 44 HCAPLUS COPYRIGHT 2002 ACS

AN 1973:77286 HCAPLUS
 DN 78:77286

TI Chromium electrodes for magnetohydrodynamic generators

IN Goolsby, Patrick F.

PA Reynolds Metals Co.

SO U.S., 4 pp.

CODEN: USXXAM

DT Patent

LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 3710152	A	19730109	US 1970-50334	19700626
	CA 967028	A1	19750506	CA 1971-116743	19710625
	GB 1358561	A	19740703	GB 1971-30222	19710628

PRAI US 1970-50334 19700626

AB Cr-based electrodes which are good thermionic **emitters** at elevated temps. are made. A Cr sample was fabricated to a typical MHD electrode configuration and exposed to temps. up to 1600.degree. in Ar for .apprx.15 hr. The wt. loss expected from vaporization did not occur. The surface was coated with a thin layer of chromic oxide, which suppressed the vapor pressure of the metal. Thus, Cr could be an ideal base electrode material. The choice of dopant materials is limited to oxides. Oxides suitable for increasing the abrasive resistance of the Cr and decreasing the thermionic work function include BaO, CaO, CeO₂, HfO₂, La₂O₃, NiO, SrO, ThO₂, TiO₂, and ZrO₂ and their mixts. The metallic compn. was formed by thoroughly mixing powd. Cr with a **metal**

oxide at 0.5-25 wt. % of the mixt. The metal oxide had thermionic emissivity and was capable of lowering the total thermionic work function of the metallic compn. relative to Cr in a carrier liq. having .apprx.3 wt. % of a surface wetter. A dried powder was formed by vacuum drying. The dried powder was pressed into buttons having the configuration of a MHD electrode at .apprx.40,000 psi and then sintered. These buttons can be used as cathodes in a MHD generator without long-term elec. degrdn., sublimative or ablative erosion, or attack by basic slags.